

## PROTOCOLADHERIN MATERIALS AND METHODS

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### FIELD OF THE INVENTION

The present invention relates, in general, to materials and methods relevant to cell-cell adhesion. More particularly, the invention relates to novel adhesion proteins, designated protocadherins, and to polynucleotide sequences encoding the protocadherins. The invention also relates to methods for inhibiting binding of the protocadherins to their natural ligands/antiligands.

### BACKGROUND

*In vivo*, intercellular adhesion plays an important role in a wide range of events including morphogenesis and organ formation, leukocyte extravasion, tumor metastasis and invasion, and the formation of cell junctions. Additionally, cell-cell adhesion is crucial for the maintenance of tissue integrity.

Intercellular adhesion is mediated by specific cell surface adhesion molecules. Cell adhesion molecules have been classified into at least four families including the immunoglobulin superfamily, the integrin superfamily, the selectin family and the cadherin superfamily. All cell types that form solid tissues express some members of the cadherin superfamily suggesting that cadherins are involved in selective adhesion of most cell types.

Cadherins have been generally described as glycosylated integral membrane proteins that have an N-terminal extracellular domain (the N-terminal 113 amino acids of the domain appear to be directly involved in binding) consisting of five subdomains characterized by sequences unique to cadherins, a hydrophobic membrane-spanning domain and a C-terminal cytoplasmic domain that interacts with the cytoskeleton through catenins and other cytoskeleton-

associated proteins. Some cadherins lack a cytoplasmic domain, however, and appear to function in cell-cell adhesion by a different mechanism than cadherins having a cytoplasmic domain. The cytoplasmic domain is required for the adhesive function of the extracellular domain in cadherins that do have an cytoplasmic domain. Binding between members of the cadherin family expressed on different cells is homophilic (*i.e.*, a member of the cadherin family binds to cadherins of its own or a closely related subclass) and  $\text{Ca}^{2+}$ -dependent. For recent reviews on cadherins, see Takeichi, *Annu. Rev. Biochem.*, 59: 237-252 (1990) and Takeichi, *Science*, 251: 1451-1455 (1991).

The first cadherins to be described (E-cadherin in mouse epithelial cells, L-CAM in avian liver, uvomorulin in the mouse blastocyst, and CAM 120/80 in human epithelial cells) were identified by their involvement in  $\text{Ca}^{2+}$ -dependent cell adhesion and their unique immunological characteristics and tissue localization. With the later immunological identification of N-cadherin, which was found to have a different tissue distribution than E-cadherin, it became apparent that a new family of  $\text{Ca}^{2+}$ -dependent cell-cell adhesion molecules had been discovered.

The molecular cloning of the genes encoding E-cadherin [see Nagafuchi *et al.*, *Nature*, 329: 341-343 (1987)], N-cadherin [Hatta *et al.*, *J. Cell. Biol.*, 106: 873-881 (1988)], and P-cadherin [Nose *et al.*, *EMBO J.*, 6: 3655-3661 (1987)] provided structural evidence that the cadherins comprised a family of cell adhesion molecules. Cloning of L-CAM [Gallin *et al.*, *Proc. Natl. Acad. Sci. USA*, 84: 2808-2812 (1987)] and uvomorulin [Ringwald *et al.*, *EMBO J.*, 6: 3647-3653 (1986)] revealed that they were identical to E-cadherin. Comparisons of the amino acid sequences of E-, N-, and P-cadherins showed a level of amino acid similarity of about 45%-58% among the three subclasses. Liaw *et al.*, *EMBO J.*, 9: 2701-2708 (1990) describes the use of PCR with degenerate oligonucleotides based on conserved regions of the E-, N- and P-cadherins to amplify N- and P-cadherin from a bovine microvascular endothelial cell cDNA.

The isolation by PCR of eight additional cadherins was reported in Suzuki *et al.*, *Cell Regulation*, 2: 261-270 (1991). Subsequently, several other cadherins were described including R-cadherin [Inuzuka *et al.*, *Neuron*, 7: 69-79 (1991)], M-cadherin [Donalies, *Proc. Natl. Acad. Sci. USA*, 88: 8024-8028 (1991)], B-cadherin [Napolitano, *J. Cell. Biol.*, 113: 893-905 (1991)] and T-cadherin [Ranscht, *Neuron*, 7: 391-402 (1991)].

Additionally, proteins distantly related to cadherins such as desmoglein [Goodwin *et al.*, *Biochem. Biophys. Res. Commun.*, 173: 1224-1230 (1990) and Koch *et al.*, *Eur. J. Cell Biol.*, 53: 1-12 (1990)] and the desmocollins [Holton *et al.*, *J. Cell Science*, 97: 239-246 (1990)] have been described. The extracellular domains of these molecules are structurally related to the extracellular domains of typical cadherins, but each has a unique cytoplasmic domain. Mahoney *et al.*, *Cell*, 67: 853-868 (1991) describes a tumor suppressor gene of *Drosophila*, called *far*, that also encodes a cadherin-related protein. The *far* tumor suppressor comprises 34 cadherin-like subdomains followed by four EGF-like repeats, a transmembrane domain, and a novel cytoplasmic domain. The identification of these cadherin-related proteins is evidence that a large superfamily characterized by a cadherin extracellular domain motif exists.

Studies of the tissue expression of the various cadherin-related proteins reveal that each subclass of molecule has a unique tissue distribution pattern. For example, E-cadherin is found in epithelial cells while N-cadherin is found in neural and muscle cells. Expression of cadherin-related proteins also appears to be spatially and temporally regulated during development because individual proteins appear to be expressed by specific cells and tissues at specific developmental stages [for review see Takeichi (1991), *supra*]. Both the ectopic expression of cadherin-related proteins and the inhibition of native expression of cadherin-related proteins hinders the formation of normal tissue structure [Detrick *et al.*, *Neuron*, 4: 493-506 (1990); Fujimori *et al.*, *Development*, 110: 97-104 (1990); Kintner, *Cell*, 69: 225-236 (1992)].

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The unique temporal and tissue expression pattern of the different cadherins and cadherin-related proteins is particularly significant when the role each subclass of proteins may play *in vivo* in normal events (e.g., the maintenance of the intestinal epithelial barrier) and in abnormal events (e.g., tumor metastasis or inflammation) is considered. Different subclasses or combinations of subclasses of cadherin-related proteins are likely to be responsible for different cell-cell adhesion events in which therapeutic detection and/or intervention may be desirable. For example, auto-antibodies from patients with pemphigus vulgaris, an autoimmune skin disease characterized by blister formation caused by loss of cell adhesion, react with a cadherin-related protein offering direct support for adhesion function of cadherins *in vivo* [Amagai *et al.*, *Cell*, 67: 869-877 (1991)]. Studies have also suggested that cadherins and cadherin-related proteins may have regulatory functions in addition to adhesive activity. Matsunaga *et al.*, *Nature*, 334: 62-64 (1988) reports that N-cadherin has neurite outgrowth promoting activity. The *Drosophila fat* tumor suppressor gene appears to regulate cell growth and suppress tumor invasion as does mammalian E-cadherin [see Mahoney *et al.*, *supra*; Frixen *et al.*, *J. Cell. Biol.*, 113:173-185 (1991); Chen *et al.*, *J. Cell. Biol.*, 114:319-327 (1991); and Vleminckx *et al.*, *Cell*, 66:107-119 (1991)]. Thus, therapeutic intervention in the regulatory activities of cadherin-related proteins expressed in specific tissues may be desirable.

There thus continues to exist a need in the art for the identification and characterization of additional cadherin-related proteins which participate in cell-cell adhesion and/or regulatory events. Moreover, to the extent that cadherin-related proteins might form the basis for the development of therapeutic and diagnostic agents, it is essential that the genes encoding the proteins be cloned. Information about the DNA sequences and amino acid sequences encoding the cadherin-related proteins would provide for the large scale production of the proteins by recombinant techniques and for the identification of the tissues/cells naturally producing the proteins. Such sequence information would also permit

the preparation of antibody substances or other novel binding molecules specifically reactive with the cadherin-related proteins that may be useful in modulating the natural ligand/antiligand binding reactions in which the proteins are involved.

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### SUMMARY OF THE INVENTION

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The present invention provides cadherin-related materials and methods that are relevant to cell-cell adhesion. In one of its aspects, the present invention provides purified and isolated polynucleotides (*e.g.*, DNA and RNA, both sense and antisense strands) encoding the novel cell adhesion molecules designated herein as protocadherins, including protocadherin-42, protocadherin-43, protocadherin pc3, protocadherin pc4 and protocadherin pc5. Preferred polynucleotide sequences of the invention include genomic and cDNA sequences as well as wholly or partially synthesized DNA sequences, and biological replicas thereof (*i.e.*, copies of the sequences made *in vitro*). Biologically active vectors comprising the polynucleotide sequences are also contemplated.

Specifically illustrating protocadherin polynucleotide sequences of the present invention are the inserts in the plasmids pRC/RSV-pc42 and pRC/RSV-pc43 which were deposited with the American Type Culture Collection (ATCC), 12301 Parklawn Drive, Rockville, Maryland 20852 on December 16, 1992 and were assigned ATCC Accession Nos. 69162 and 69163, respectively.

The scientific value of the information contributed through the disclosures of the DNA and amino acid sequences of the present invention is manifest. For example, knowledge of the sequence of a partial or complete DNA encoding a protocadherin makes possible the isolation by standard DNA/DNA hybridization or PCR techniques of full length cDNA or genomic DNA sequences that encode the protein (or variants thereof) and, in the case of genomic DNA sequences, that specify protocadherin-specific regulatory sequences such as promoters, enhancers and the like. Alternatively, DNA sequences of the present invention may be chemically synthesized by conventional techniques.

Hybridization and PCR techniques also allow the isolation of DNAs encoding heterologous species proteins homologous to the protocadherins specifically illustrated herein.

5 According to another aspect of the invention, host cells, especially eucaryotic and procaryotic cells, are stably transformed or transfected with the polynucleotide sequences of the invention in a manner allowing the expression of protocadherin polypeptides in the cells. Host cells expressing protocadherin polypeptide products, when grown in a suitable culture medium, are particularly useful for the large scale production of protocadherin polypeptides, fragments and variants thereby enabling the isolation of the desired polypeptide products from the cells or from the medium in which the cells are grown.

10 The novel protocadherin protein products of the invention may be obtained as isolates from natural tissue sources, but are preferably produced by recombinant procedures involving the host cells of the invention. The products may be obtained in fully or partially glycosylated, partially or wholly de-glycosylated, or non-glycosylated forms depending on the host cell selected or recombinant production and/or post-isolation processing.

15 Protocadherin variants according to the invention may comprise polypeptide analogs wherein one or more of the specified amino acids is deleted or replaced or wherein one or more non-naturally encoded amino acids are added: 20 (1) without loss, and preferably with enhancement, of one or more of the biological activities or immunological characteristics specific for a protocadherin; or (2) with specific disablement of a particular ligand/antiligand binding function. Also contemplated by the present invention are antibody substances (e.g., 25 monoclonal and polyclonal antibodies, chimeric and humanized antibodies, antibody domains including Fab, Fab', F(ab')<sub>2</sub>, Fv or single variable domains, and single chain antibodies) which are specific for the protocadherins of the invention. Antibody substances can be developed using isolated natural, recombinant or synthetic protocadherin polypeptide products or host cells

expressing such products on their surfaces. The antibody substances may be utilized for purifying protocadherin polypeptides of the invention, for determining tissue expression of polypeptides and as antagonists of the ligand/antiligand binding activities of the protocadherins. Specifically illustrating monoclonal antibodies of the present invention are the protocadherin-43 specific monoclonal antibodies produced by the hybridoma cell line designated 38I2C which was deposited with the ATCC on December 2, 1992 and was assigned ATCC Accession No. HB 11207.

Numerous other aspects and advantages of the present invention will be apparent upon consideration of the following detailed description, reference being made to the drawing wherein FIGURE 1A-C is an alignment of protocadherin amino acid sequences of the invention with the amino acid sequences of N-cadherin and of the *Drosophila fat* tumor suppressor.

#### DETAILED DESCRIPTION

The present invention is illustrated by the following examples wherein Examples 1, 2 and 3 describe the isolation by PCR of protocadherin polynucleotide sequences. Example 3 also describes the chromosome localization of several protocadherin genes of the invention. Example 4 describes the isolation by DNA/DNA hybridization of additional protocadherin polynucleotide sequences of the present invention. Example 5 presents the construction of expression plasmids including polynucleotides encoding protocadherin-42 or protocadherin-43 and the transfection of L cells with the plasmids. The generation of antibodies to protocadherin-42 and protocadherin-43 is described in Example 6. Example 7 presents the results of immunoassays of transfected L cells for the expression of protocadherin-42 or protocadherin-43. Example 8 describes the cell aggregation properties of L cells transfected with protocadherin-42, protocadherin-43 or a chimeric protocadherin-43/E-cadherin molecule. The calcium-binding properties of pc43 are described in Example 9. The results of assays of various tissues and cell lines for the expression of protocadherin-42 and protocadherin-43

by Northern blot, Western blot and *in situ* hybridization are respectively presented in Examples 10, 11 and 12. Example 13 describes immunoprecipitation experiments identifying a 120 kDa protein that coprecipitates with protocadherin-43.

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### Example 1

The polymerase chain reaction (PCR) was used to isolate novel rat cDNA fragments encoding cadherin-related polypeptides.

#### Design of PCR Primers

10 Two regions of conserved amino acid sequence, one from the middle of the third cadherin extracellular subdomain (EC-3) and the other from the C-terminus of the fourth extracellular subdomain (EC-4), were identified by comparison of the published amino acid sequences for L-CAM (Gallin *et al.*, *supra*), E-cadherin (Nagafuchi *et al.*, *supra*), mouse P-cadherin (Nose *et al.*, *supra*), uvomorulin (Ringwald *et al.*, *supra*), chicken N-cadherin (Hatta *et al.*, *supra*), mouse N-cadherin [Miyatani *et al.*, *Science*, 245:631-635 (1989)] and human P-cadherin [Shimoyama *et al.*, *J. Cell. Biol.*, 109:1787-1794 (1989)], and the corresponding degenerate oligonucleotides respectively set out below in IUPAC-IUB Biochemical nomenclature were designed for use as PCR primers.

20 Primer 1 (SEQ ID NO: 1)  
5' AARSSNNTNGAYTRYGA 3'  
Primer 2 (SEQ ID NO: 2)  
3' TTRCTRTTRCGNGGNNN 5'

The degenerate oligonucleotides were synthesized using an Applied Biosystems model 380B DNA synthesizer (Foster City, California).

#### Cloning of cDNA Sequences by PCR

25 PCR was carried out in a manner similar to that described in Suzuki *et al.*, *Cell Regulation*, 2: 261-270 (1991) on a rat brain cDNA preparation. Total RNA was prepared from rat brain by the guanidium

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isothiocyanate/cesium chloride method described in Maniatis *et al.*, pp. 196 in *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor, New York: Cold Spring Harbor Laboratory (1982). Brain poly(A)<sup>+</sup> RNAs were then isolated using a FastTrack<sup>®</sup> kit (Invitrogen, San Diego, California) and cDNA was prepared using a cDNA synthesis kit (Boehringer Mannheim Biochemicals, Indianapolis, Indiana). The PCR reaction was initiated by adding 2.5 units of Taq DNA polymerase (Boehringer Mannheim Biochemicals) to 100 ng template cDNA and 10 µg of each primer, after which 35 reaction cycles of denaturation at 94°C for 1.5 minutes, annealing at 45°C for 2 minutes, and polymerization at 72°C for 3 minutes were carried out. Two major bands of about 450 base pairs (bp) and 130 bp in size were found when the products of the PCR reaction were subjected to agarose gel electrophoresis. The 450 bp band corresponded to the expected length between the two primer sites corresponding to the middle of the third cadherin extracellular subdomain (EC-3) and the carboxyl terminus of the fourth cadherin extracellular subdomain (EC-4), but the 130 bp band could not be predicted from any of the previously identified cadherin sequences. The 450 bp and 130 bp bands were extracted by a freezing and thawing method. The resulting fragments were phosphorylated at the 5' end with T4 polynucleotide kinase and subcloned by a blunt-end ligation into the Sma I site of M13mp18 (Boehringer Mannheim Biochemicals) in a blunt end ligation for sequence analysis. Sequencing of the fragments was carried out by the dideoxynucleotide chain termination method using a Sequenase kit (United States Biochemicals, Cleveland, Ohio). DNA and amino acid sequence were analyzed using the Beckman Microgenie program (Fullerton, California).

#### Analysis of cDNA Sequences

Nineteen novel partial cDNA clones were isolated. The DNA and deduced amino acid sequences of the clones (including sequences corresponding to the PCR primers) are set out as follows: RAT-123 (SEQ ID NOs: 3 and 4, respectively), RAT-212 (SEQ ID NOs: 5 and 6), RAT-214 (SEQ ID NOs: 7 and

8), RAT-216 (SEQ ID NOs: 9 and 10), RAT-218 (SEQ ID NOs: 11 and 12), RAT-224 (SEQ ID NOs: 13 and 14), RAT-312 (SEQ ID NOs: 15 and 16), RAT-313 (SEQ ID NOs: 17 and 18), RAT-314 (SEQ ID NOs: 19 and 20), RAT-315 (SEQ ID NOs: 21 and 22), RAT-316 (SEQ ID NOs: 23 and 24), RAT-317 (SEQ ID NOs: 25 and 26), RAT-321 (SEQ ID NOs: 27 and 28), RAT-323 (SEQ ID NOs: 29 and 30), RAT-336 (SEQ ID NOs: 31 and 32), RAT-352 (SEQ ID NOs: 33 and 34), RAT-411 (SEQ ID NOs: 35 and 36), RAT-413 (SEQ ID NOs: 37 and 38), and RAT-551 (SEQ ID NOs: 39 and 40).

The deduced amino acid sequences of the cDNA clones are homologous to, but distinct from the known cadherins. The cadherins described thus far have highly conserved, short amino acid sequences in the third extracellular subdomain (EC-3) including the consensus sequence D-Y-E or D-F-E located at the middle region of the subdomain and the consensus sequence D-X-N-E-X-P-X-F (SEQ ID NO: 41) or D-X-D-E-X-P-X-F (SEQ ID NO: 42) at its end (Hatta et al., *supra*), while the corresponding sequences of other subdomains, except for the fifth extracellular subdomain (EC-5), are D-R-E and D-X-N-D-N-X-P-X-F (SEQ ID NO: 43), respectively. In contrast, the deduced amino acid sequences of the new clones that correspond to cadherin extracellular subdomains include the sequence D-Y-E or D-F-E at one end, but have the sequence D-X-N-D-N-X-P-X-F instead of D-X-N-E-X-P-X-F or D-X-D-E-X-P-X-F, at the other end. The polypeptides encoded by the partial clones are homologous to previously identified cadherins but did not show significant homology to any other sequences in Genbank. Therefore, the partial cDNAs appear to comprise a new subclass of cadherin-related molecules.

### Example 2

Various cDNA fragments structurally similar to the rat cDNAs described in Example 1 were isolated from human, mouse, and Xenopus brain cDNA preparations and from *Drosophila* and *C. elegans* whole body cDNA

preparations by PCR using Primers 1 and 2 as described in Example 1. The DNA and deduced amino acid sequences of the resulting PCR fragments (including sequences corresponding to the PCR primers) are set out as follows: MOUSE-321 (SEQ ID NOs: 44 and 45), MOUSE-322 (SEQ ID NOs: 46 and 47),  
5      MOUSE-324 (SEQ ID NOs: 48 and 49), MOUSE-326 (SEQ ID NOs: 50 and 51),  
HUMAN-11 (SEQ ID NOs: 52 and 53), HUMAN-13 (SEQ ID NOs: 54 and 55),  
HUMAN-21 (SEQ ID NOs: 56 and 57), HUMAN-24 (SEQ ID NOs: 58 and 59),  
HUMAN-32 (SEQ ID NOs: 60 and 61), HUMAN-42 (SEQ ID NOs: 62 and 63),  
HUMAN-43 (SEQ ID NOs: 64 and 65), HUMAN-212 (SEQ ID NOs: 66 and  
10      67), HUMAN-213 (SEQ ID NOs: 68 and 69), HUMAN-215 (SEQ ID NOs: 70  
and 71), HUMAN-223 (SEQ ID NOs: 72 and 73), HUMAN-410 (SEQ ID NOs:  
74 and 75), HUMAN-443 (SEQ ID NOs: 76 and 77), XENOPUS-21 (SEQ ID  
NOs: 78 and 79), XENOPUS-23 (SEQ ID NOs: 80 and 81), XENOPUS-25 (SEQ  
ID NOs: 82 and 83), XENOPUS-31 (SEQ ID NOs: 84 and 85), DROSOPHILA-  
15      12 (SEQ ID NOs: 86 and 87), DROSOPHILA-13 (SEQ ID NOs: 88 and 89),  
DROSOPHILA-14 (SEQ ID NOs: 90 and 91) and C.ELEGANS-41 (SEQ ID  
NOs: 92 and 93). Comparison of the deduced amino acid sequences indicates  
significant similarity between sets of these clones. In particular, there are three  
sets of clones that appear to be cross-species homologues: RAT-218, MOUSE-322  
20      and HUMAN-43; RAT-314, MOUSE-321 and HUMAN-11; and MOUSE-326  
and HUMAN-42.

### Example 3

To ascertain the complete structure of the new proteins defined by the PCR products, two full length human cDNAs corresponding to the partial  
25      cDNAs HUMAN-42 and HUMAN-43 were isolated.

#### Isolation of Full-length Human cDNAs

A human fetal brain cDNA library (Stratagene, La Jolla, California) in the  $\lambda$ ZapII vector was screened by the plaque hybridization method

[described in Ausubel *et al.*, Eds., *Current Protocols in Molecular Biology*, Sections 6.1.1 to 6.1.4 and 6.2.1 to 6.2.3, John Wiley & Sons, New York (1987)] with <sup>32</sup>P-labelled HUMAN-42 and HUMAN-43 DNA fragments. The positive clones were plaque-purified and, using a helper virus, the inserts were cut out by an *in vivo* excision method in the form of a Bluescript SK(+) plasmid. The insert sequences were then subcloned into the M13 vector (Boehringer Mannheim, Biochemicals) for sequencing. Several overlapping cDNA clones were isolated with each probe including two cDNAs which contained the putative entire coding sequences of two novel proteins designated protocadherin-42 (pc42) and protocadherin-43 (pc43). The DNA and deduced amino acid sequences of pc42 are set out in SEQ ID NOs: 94 and 95, respectively, while the DNA and deduced amino acid sequences of pc43 are set out in SEQ ID NOs: 96 and 97, respectively.

A description of the cloning of protocadherin sequences of the invention was published in Sano *et al.*, *The EMBO Journal*, 12(6): 2249-2256 (1993) after filing of the priority application hereto. The deduced amino acid sequence of pc43 was previously presented at the December 9, 1991 meeting of the American Society for Cell Biology. An abstract of the presentation is published as Suzuki *et al.*, *J. Cell. Biol.*, 115: 72a (Abstract 416) (December 9, 1991).

#### Analysis of Full-length Human Clones

Comparison of the full length cDNA sequences of pc42 and pc43 to the sequences of the various DNA fragments originally obtained by PCR reveals that MOUSE-326 and HUMAN-42 correspond to a portion of the fourth extracellular subdomain (EC-4) of pc42, and RAT-314, MOUSE-321, and HUMAN-11 correspond to a portion of the third extracellular subdomain (EC-3) of pc43 and RAT-218, MOUSE-322 and HUMAN-43 correspond to a portion of the fifth extracellular domain (EC-5) of pc43.

5 The overall structures of pc42 and pc43 are similar to that of  
typical cadherins but the new molecules also have distinct features. Both  
protocadherin cDNA sequences contain putative translation initiation sites and  
translated amino acid sequences start with typical signal sequences, but the clones  
10 lack the prosequences that are present in all known cadherin precursors. The  
cDNAs encode proteins having a large N-terminal extracellular domain and a  
relatively short C-terminal cytoplasmic domain connected by a transmembrane  
sequence. The extracellular domains of pc42 and pc43 are different in length and  
pc42 contains seven subdomains that closely resemble the typical cadherin  
15 extracellular subdomain while pc43 has six such subdomains. The sizes of the  
protocadherin cytoplasmic domains are similar to those of typical cadherins, but  
the sequences do not show any significant homology with those of known  
cadherins or cadherin-related proteins.

20 Amino acid identity determinations between extracellular  
subdomains of human pc42 and pc43, and of mouse N-cadherin (SEQ ID NO: 98)  
(presented as an example of a "typical" cadherin) and the eighteenth extracellular  
subdomain of *Drosophila fat* tumor suppressor (EC-18, SEQ ID NO: 99) (the  
eighteenth extracellular subdomain of *fat* is a prototypical *fat* subdomain) are  
presented in Table 1 below, wherein, for example, "N-EC-1 x pc42" indicates  
that the first extracellular subdomain of N-cadherin was compared to the  
extracellular subdomain of pc42 indicated on the horizontal axis.

Table 1

		<u>EC-1</u>	<u>EC-2</u>	<u>EC-3</u>	<u>EC-4</u>	<u>EC-5</u>	<u>EC-6</u>	<u>EC-7</u>
5	N-EC-1 x pc42	20	27	26	26	31	29	17
	N-EC-1 x pc43	31	23	23	26	31	24	
	N-EC-2 x pc42	28	30	32	30	37	31	19
	N-EC-2 x pc43	30	28	30	36	29	30	
	N-EC-3 x pc42	21	26	30	29	31	30	22
10	N-EC-3 x pc43	25	18	26	28	28	25	
	N-EC-4 x pc42	28	28	26	25	29	27	17
	N-EC-4 x pc43	21	25	28	28	29	24	
	N-EC-5 x pc42	24	21	25	24	24	19	12
	N-EC-5 x pc43	15	21	20	20	25	16	
	fat EC-18 x pc42	22	35	32	34	42	35	19
	fat EC-18 x pc43	32	30	36	36	33	29	

15 The amino acid identity values between the extracellular subdomains of pc42 and pc43, and N-cadherin EC-1 through EC-5 and *Drosophila fat* EC-18 are mostly less than 40%. These identity values are comparable to the values between the subdomains of other cadherin subclasses. However, higher identity values indicate that pc42 and pc43 are more closely related to *fat* than to N-cadherin.

20 Amino acid identity determinations between extracellular subdomains of human pc42 and pc43 are presented in Table 2 below.

Table 2

	pc42						
<u>pc43</u>	<u>EC-1</u>	<u>EC-2</u>	<u>EC-3</u>	<u>EC-4</u>	<u>EC-5</u>	<u>EC-6</u>	<u>EC-7</u>
EC-1	33	27	29	26	25	26	25
EC-2	26	38	29	33	34	28	21
EC-3	26	32	41	30	32	31	22
EC-4	25	34	30	41	39	31	18
EC-5	23	32	29	27	36	34	16
EC-6	25	25	26	25	28	23	26

The identity values between respective EC-1, EC-2, EC-3, EC-4, EC-5 subdomains and the last subdomains of pc42 and pc43 are generally higher values than values obtained for comparisons of the protocadherins to N-cadherin. These results suggest that pc42 and pc43 are more closely related to one another than they are to classic cadherins.

FIGURE 1A-C presents an alignment of the deduced amino acid sequences of the extracellular subdomains of pc42 (EC-1 through EC-7), pc43 (EC-1 through EC-6), mouse N-cadherin (EC-1 through EC-5) and *Drosophila fat* EC-18. A sequence on a line in FIGURE 1A continues on the same line in FIGURES 1B and 1C. Gaps were introduced to maximize homology. The amino acid residues described by capital letters in the "motif" line are present in more than half of the subdomains of N-cadherin, pc42, pc43 and *Drosophila fat*. The amino acid residues described by small letters in the motif line are less well conserved in human pc42, pc43, and *Drosophila fat*. FIGURE 1A-C shows that many amino acids characteristic of other cadherin extracellular domain repeats are conserved in the pc42 and pc43 sequences, including the cadherin sequence motifs DXD, DRE and DXNDNXPXF (SEQ ID NO: 43), two glycine residues, and one glutamic acid residue. Additionally, pc42 and pc43 share unique features in comparison to N-cadherin. More amino acids at specific sites are conserved

between pc42 and pc43, such as the DXDXGXN (SEQ ID NO: 100) protocadherin sequence motif near the amino terminus of the pc42 and pc43 subdomains and the AXDXGXP (SEQ ID NO: 101) sequence motif near the carboxyl terminus of the subdomains. Additionally, both protocadherins share regions that do not show significant homology with the typical cadherin motif (of N-cadherin) near the carboxyl terminus of EC-1, in the middle of EC-2 and EC-4, and at the carboxyl terminus of the last repeat. A cysteine residue is located at a similar position in the middle of EC-4 of pc42 and pc43. In general, the extracellular subdomains of pc42 and pc43 are more similar to EC-18 of *far* than the extracellular subdomains of N-cadherin.

#### Possible Alternative Splicing

Sequence analysis of various overlapping protocadherin cDNA clones revealed that some clones contained unique sequences at the 3' end, although the 5' end sequences were identical to other clones. The sequences forming the boundaries of the 3' end regions are consistent with the consensus sequence of mRNA splicing, suggesting that these clones may correspond to alternatively spliced mRNAs. The DNA and deduced amino acid sequences of one possible product of alternative splicing of pc42 mRNA are set out in SEQ ID NOs: 102 and 103. The DNA and deduced amino acid sequences of two possible products of alternative splicing of pc43 mRNA are respectively presented in SEQ ID NO: 104 and 105, and SEQ ID NOs: 106 and 107.

#### Chromosome Localization

The chromosomal location of the protocadherin 413 gene (SEQ ID NO: 37) and of the pc42 and pc43 genes was determined by conventional methods.

Briefly, C3H/HeJ-*gld* and *Mus spretus* (Spain) mice and [(C3H/HeJ-*gld* x *Mus spretus*) F<sub>1</sub> x C3H/HeJ-*gld*] interspecies backcross mice were bred and maintained as previously described in Seldin, *et al.*, *J. Exp. Med.*, 167: 688-693 (1988). *Mus spretus* was chosen as the second parent in the cross



because of the relative ease of detection of informative restriction fragment length variants (RFLVs) in comparison with crosses using conventional inbred laboratory strains. Gene linkage was determined by segregation analysis.

5 Genomic DNA isolated from mouse organs by standard techniques  
was digested with restriction endonucleases and 10 $\mu$ g samples were  
electrophoresed in 0.9% agarose gels. DNA was transferred to Nytran  
membranes (Schleicher & Schull, Inc., Keene, NH), hybridized with the  
appropriate probe at 65°C and washed under stringent conditions, all as  
10 previously described in Maniatis *et al.*, *supra*). To localize the pc42 gene, a  
mouse sequence probe corresponding to nucleotides 1419 to 1906 of SEQ ID NO:  
94 was used and for pc43 a rat sequence probe corresponding to nucleotides 1060  
to 1811 of SEQ ID NO: 96 was used. To localize the procadherin 413 gene, a  
probe including the sequence set out in SEQ ID NO: 37 was used. Other clones  
15 used as probes in the current study and RFLVs used to detect anonymous DNA  
loci were all previously described [Chromosome 7, DNA segment, Washington  
12 (*D7Was12*); the parathyroid hormone (*Pth*); calcitonin (*Calc*); hemoglobin,  $\beta$   
chain (*Hbb*); metallothionein-I (*Mt-1*); adenine phosphoribosyltransferase (*Aprt*);  
growth hormone receptor (*Ghr*); prostaglandin E receptor EP2 subtype  
(*Ptgerp2*); dihydrofolate reductase-2 (*Dhfr2*); fibroblast growth factor a (*Fgfa*);  
20 and glucocorticoid receptor-1 (*Grl-1*)].

Comparison of the haplotype distribution of protocadherin genes  
with those determined for loci throughout the mouse genome allowed each to be  
mapped to specific regions of mouse chromosomes. The probability for linkage  
was >99% and indicated assignment of both the pc42 gene and the pc43 gene  
25 was chromosome 18. The assignment of the protocadherin 413 gene was  
chromosome 7. The region of chromosome 18 to which the pc42 and pc43 genes  
were mapped corresponds to the ataxia (*ax*) loci [Burt, *Anat. Rec.*, 196: 61-69  
(1980) and Lyon, *J. Hered.*, 46: 77-80 (1955)] and twirler (*Tw*) loci [Lyon, *J.*  
*Embryol. Exp. Morphol.*, 6: 105-116 (1958)], while the region of chromosome

7 to which the protocadherin 413 gene was mapped corresponds to the shaker (*sh-1*) locus [Kikuchi *et al.*, *Acta Oto-Laryngol.*, 60: 287-303 (1965) and Lord *et al.*, *Am. Nat.*, 63: 453-442 (1929)]. These loci have been implicated as involved in hereditary neural disease in the mouse. This result is consistent with *in situ* hybridization results (see Example 12) showing that pc42 and pc43 are strongly expressed in the brain and particularly in the cerebellum.

#### Example 4

Two additional novel human protocadherin cDNAs and one additional novel rat protocadherin cDNA were isolated using rat protocadherin fragments described in Example 1 as probes.

Initially, the rat clone RAT-214 (SEQ ID NO: 7) was used as a probe to screen a rat brain cDNA library (Stratagene, La Jolla, CA). The final washing step was performed twice at 50°C in 0.1X SSC with 0.1% SDS for 15 minutes. Various clones were identified which contained partial cDNA inserts encoding related protocadherin amino acid sequences. The nucleotide sequence of one novel rat clone designated #6-2 is set out in SEQ ID NO: 108. The first fifteen nucleotides of SEQ ID NO: 108 are the sequence of a linker and are not part of the rat #6-2 clone.

A human fetal brain cDNA library obtained from Stratagene was screened with the 0.7 kbp PstI fragment of clone #6-2. The fragment appears to encode the EC-2 and EC-3 of the rat protocadherin. After screening about  $2 \times 10^6$  phages, eleven positive clones were isolated. Sequencing of the clones identified a novel full length human protocadherin cDNA designated human pc3. The nucleotide and deduced amino acid sequence of human pc3 are set out in SEQ ID NOs: 109 and 110.

The 0.7 kbp PstI fragment of rat clone #6-2 was also used to rescreen the Stratagene rat brain cDNA library for full length rat cDNA clones. A clone containing an insert encoding a full length novel protocadherin cDNA

was isolated. The DNA and deduced amino acid sequence of the insert are set out in SEQ ID NO: 111 and 112. The full length rat cDNA was named pc5 because it does not appear to be the homolog of the human pc3 clone based upon a comparison of the sequences.

5 Concurrently, the 0.8 kbp Eco RI-Pst I fragment of partial rat cDNA designated #43 (SEQ ID NO: 113), which was obtained by screening the Stratagene rat brain cDNA library with a probe corresponding to the human pc43 cytoplasmic domain, was used to probe the Stratagene human cDNA library for full length human protocadherin cDNAs. The fragment appears to encode EC-3  
10 through the beginning of EC-6 of clone #43. One partial clone identified encodes a novel human protocadherin named human pc4. The nucleotide sequence and deduced amino acid sequences of the human pc4 clone are set out in SEQ ID NOs: 114 and 115. The amino acid sequence encoded by the pc4 clone appears to begin in the middle of EC-2 of pc4 and continues through the cytoplasmic tail  
15 of the protocadherin.

#### Example 5

The full length human cDNAs encoding pc42 and pc43 were expressed in L cells (ATCC CCL 1) using the pRC/RSV expression vector (Invitrogen, San Diego, California). The cDNAs were isolated from the  
20 Bluescript SK(+) clones described in Example 2 by digestion with SspI followed by blunt-ending with DNA polymerase and digestion with XbaI (for pc42), or by double digestion with SpeI and EcoRV (for pc43). The pRC/RSV expression vector was digested with HindIII, followed by blunt-ending and re-digestion with XbaI for insertion of pc42 sequences, or by digested with XbaI followed by  
25 blunt-ending and re-digestion with SpeI for insertion of pc43 sequences. The isolated protocadherin DNAs were ligated into the linearized pRC/RSV vector. The resulting pc42 expression plasmid designated pRC/RSV-pc42 (ATCC 69162) and pc43 expression plasmid designated pRC/RSV-pc43 (ATCC 69163) were

purified by CsCl gradient centrifugation and transfected into L cells by a Calcium phosphate method.

5 The pc42 and pc43 transfectants were morphologically similar to the parental cells. Northern blot analysis of L cells transfected with pc42 or pc43 DNA sequences showed that the transfected cells expressed mRNAs of a size expected to encode the particular protocadherin.

### Example 6

Rabbit polyclonal antibodies specific for pc42 and pc43 were generated as well as a mouse monoclonal antibody specific for pc43.

#### 10 Preparation of Polyclonal Antibodies Specific for pc42 and pc43

15 DNA sequences encoding portions of the extracellular domain of pc42 and pc43 were each fused to a maltose binding protein-encoding sequence and expressed in bacteria. Specifically, DNAs corresponding to EC-4 through EC-7 of pc42 and EC-3 through EC-5 of pc43 were prepared by PCR and subcloned in the correct reading frame into the multicloning site of the pMAL expression vector (New England Biolabs, Beverly, Massachusetts) which contains sequences encoding maltose binding protein immediately upstream of the multicloning site. The resulting plasmids were then introduced into *E. coli* NM522 cells (Invitrogen, San Diego, California) by a single step transformation method. Expression of the fusion proteins was induced by the addition of IPTG and the fusion proteins were purified from cell extracts by amylose resin affinity chromatography (New England Biolabs) as described by the manufacturer. The fusion proteins were used for the immunization of rabbits without further purification.

25 Polyclonal antibodies were prepared in rabbits by immunization at four subcutaneous sites with 500 $\mu$ g of purified fusion protein in Freund's complete adjuvant. Subsequent immunizations with 100 $\mu$ g of the fusion protein were in Freund's incomplete adjuvant. Immune sera was passed through

sepharose coupled to maltose binding protein (New England Biolabs) and polyclonal antibodies were purified from immune sera using Sepharose affinity columns prepared by reaction of the purified fusion protein with CNBr Sepharose (Pharmacia). Reactivity of the polyclonal sera with purified pc42 fusion protein and pc42 transfected cell extracts (described in Example 5) was confirmed.

#### Preparation of Monoclonal Antibodies Specific for pc43

The pc43 fusion protein (containing the EC-3 through EC-5 subdomains of pc43) was used to generate monoclonal antibodies in mice according to the method of Kennett, *Methods in Enzymol.*, 58:345-359 (1978). Briefly, mice were immunized with the pc43 fusion protein (100 $\mu$ g) at two subcutaneous sites. The spleen from the highest titer mouse was fused to the NS1 myeloma cell line. The resulting hybridoma supernatants were screened in a ELISA assay for reactivity with the pc43 fusion protein and with maltose binding protein. The fusion wells with the highest reactivity to the pc43 extracellular domains were subcloned. The hybridoma cell line designated 38I2C (ATCC HB 11207) produced a IgG<sub>1</sub> subtype monoclonal antibody specific for pc43. Reactivity of the monoclonal antibody produced by hybridoma cell line 38I2C to pc43 was confirmed by immunoblotting the pc43 L cell transfectants described in Example 5. The 38I2C monoclonal antibody is specific for human pc43.

#### Example 7

L cells transfected with DNA sequences encoding pc42 and pc43 as prepared in Example 5 were assayed for expression of the protocadherins by immunoblot and by immunofluorescence microscopy.

#### Immunoblot Analysis

Cell extracts of pc42 and pc43 transfectants were subjected to SDS-PAGE and then blotted electrophoretically onto a PVDF membrane (Millipore, Bedford, Massachusetts). The membranes were incubated with 5% skim milk in Tris-buffered saline (TBS) for two hours and then respectively with

either pc42 polyclonal sera or pc43 monoclonal antibody for one hour. The membranes were washed three times (for 5 minutes each wash) with TBS containing 0.05% Tween 20 and respectively incubated with alkaline phosphatase-conjugated anti-rabbit IgG antibody or anti-mouse IgG antibody (Promega, Madison, Wisconsin) in the same buffer for one hour. After washing the membranes with TBS containing 0.05% Tween 20, reactive bands were visualized by using Western Blue solution (Promega).

Anti-pc42 polyclonal antibodies stained a band of about 170 kDa molecular weight in pc42 transfected cells, but not parental L cells. The pc43-specific monoclonal antibody (38I2C) and polyclonal antibodies stained two adjacent bands of about 150 kDa molecular weight in pc43 transfected cells. The pc43 antibodies did not stain bands in parental L-cells. The molecular weights indicated by the staining of bands by the pc42 and pc43 antibodies are significantly larger than the molecular weights predicted from the deduced amino acid sequences. This discrepancy in molecular weight is common among various cadherin-related proteins and may be attributable to the glycosylation and/or cadherin specific structural properties. The pc42 antibody also stained smaller bands, which may be proteolytic degradation products.

When transfected cells were trypsinized and cell extracts were prepared, run on SDS/PAGE and immunoblotted with the appropriate antibody, the pc42 and pc43 polypeptides expressed by the transfected cells were found to be highly sensitive to proteolysis and were easily digested by 0.01% trypsin treatment. In contrast to the classic cadherins, however, these proteins were not protected from the digestion in the presence of 1-5mM  $\text{Ca}^{2+}$ .

#### Immunofluorescence Microscopy

Transfected cells were grown on a cover slip precoated with fibronectin and were fixed with 4% paraformaldehyde for 5 minutes at room temperature or with cold methanol on ice for 10 minutes followed by 4% paraformaldehyde fixation. After washing with TBS, the cells were incubated with

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5 TBS containing 1% BSA for 30 minutes and then with anti-pc42 polyclonal antibody or anti-pc43 monoclonal antibody in TBS containing 1% BSA for 1 hour at room temperature. Cover slips were then washed with TBS containing 0.01% BSA and respectively incubated with FITC-conjugated anti-rabbit antibody or anti-mouse antibody (Cappel, Durham, North Carolina) for 60 minutes at room temperature. The cells were washed again with TBS containing 0.01% BSA and subjected to fluorescence microscopy. Both pc42-specific and pc43-specific polyclonal antibodies stained the cell periphery of transfected cells expressing the protocadherin proteins, mainly at the cell-cell contact sites. The antibodies did not stain the parent L cells, nor did rabbit preimmune sera stain the pc42 and pc43 transfectants.

#### Example 8

15 The cell aggregation properties of the transfected L cells expressing protocadherin proteins were examined. Transfected L cells were cultured in Dulbecco's Modified Eagles Medium (DMEM) (Gibco, Grand Island, New York) supplemented with 10% fetal bovine serum at 37°C in 5% CO<sub>2</sub>. Cells grown near confluence were treated with 0.01% trypsin in the presence of 1 mM EGTA for 25 minutes on a rotary shaker at 37°C and collected by centrifugation. The cells were washed three times with Ca<sup>2+</sup> free HEPES-buffered saline (HBS) after adding soybean trypsin inhibitor, and were resuspended in HBS containing 1% BSA. The cell aggregation assay [Urushihara *et al.*, *Dev. Biol.*, 70: 206-216 (1979)] was performed by incubating the resuspended cells in a 1:1 mixture of DMEM and HBS containing 1% BSA, 2 mM CaCl<sub>2</sub> and 20 µg/ml of deoxyribonuclease on a rotary shaker at 37°C for 30 minutes to 6 hours.

25 The pc42 and pc43 transfectants did not show any significant cell aggregation activity during periods of incubation less than 1 hour. This is in contrast to the cell aggregation that occurs with classic cadherins in similar experiments (Nagafuchi *et al.*, *supra*, and Hatta *et al.*, *supra*). However,

prolonged incubation of transfected cells (more than 1-2 hours) resulted in gradual re-aggregation of the cells into small aggregates. Similar results were obtained when single cell suspensions of transfected cells were prepared by trypsin treatment in the presence of  $\text{Ca}^{2+}$ . No re-aggregation was observed under the same conditions when untransfected L cells or L cells transfected with pRC/RSV vector alone were tested. When pc43 transfectants labelled with DiO (Molecular Probes, Eugene, OR) were incubated with unlabelled pc42 transfectants in the cell aggregation assay, aggregation of labelled and unlabelled cells was almost mutually exclusive indicating that protocadherin binding is homophilic.

In view of the fact that the protocadherin cytoplasmic domains exhibit no apparent homology to cadherin domains, experiments were performed to determine if the difference in cytoplasmic domains could account for the difference in cell aggregation activity observed in cadherin and protocadherin transfectants. The cytoplasmic domain of pc43 was replaced with the cytoplasmic domain of E-cadherin and aggregation of cells transfected with the chimeric construct was analyzed.

The Bluescript SK(+) clone described in Example 2 which contained the entire coding sequence for pc43 was digested with EcoRV and then partially digested with XbaI to remove the sequence corresponding to the cytoplasmic domain, and the plasmid DNA was purified by agarose gel electrophoresis. The cDNA corresponding to the cytoplasmic domain of mouse E-cadherin was synthesized by PCR using mouse cDNA made from mouse lung mRNA as a template and specific primers corresponding to a region near the N-terminus of the cytoplasmic domain sequence or the region containing the stop codon of mouse E-cadherin (Nagafuchi *et al.*, *supra*). A XbaI sequence was included to the 5' end of the upstream primer. The E-cadherin cytoplasmic domain cDNA was then subcloned into the linearized pc43 Bluescript clone. The DNA containing the entire resulting chimeric sequence was cut out with SpeI and EcoRV and was subcloned into the SpeI-blunted XbaI site of the expression vector pRc/RSV vector. Finally, L cells were transfected with the resultant construct by



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5 a calcium phosphate method. After screening with G418 for about 10 days, the transfectants were stained with FITC-labeled 38I2C anti-pc43 antibody and subjected to FACS analysis. A portion of highly labeled cells were isolated and cloned. Transfectants showed a morphology similar to that of parental L cells and the expressed protein was localized at the cell periphery using pc43 antibody for immunofluorescence microscopy.

10 Cell aggregation activity of the chimeric transfectants was analyzed as follows. The chimeric pc43 transfectants were labeled with DiO for 20 minutes at room temperature. The resultant cells were trypsinized in the presence of 1mM EGTA and single cell suspension was made. Then, the cells were mixed with unlabeled other type of transfectants and incubated on a rotary shaker for two hours. The results were examined with a fluorescence and a phase contrast microscope apparatus. Antibody inhibition of cell aggregation was examined by incubation of the transfectants in the presence of polyclonal anti-pc43 antibody

15 (100 ng/ml) in the standard assay medium.

In the cell aggregation assay, the chimeric pc43 transfectants showed clear  $\text{Ca}^{2+}$ -dependent cell aggregation within forty minutes of incubation. Cell aggregation was inhibited by the addition of pc43-specific polyclonal antibody.

#### 20 Example 9

25 The procedures of Maruyama *et al.*, *J. Biochem.*, 95: 511-519 (1984) were used to determine the calcium binding properties of pc43 by Western blot analysis in the presence or absence of calcium-45. The pc43 fusion protein described in Example 6 containing pc43 subdomains EC-3 through EC-5 was compared to the calcium binding protein calmodulin. Samples of purified pc43 fusion protein were run on SDS/PAGE and electrophoretically transferred to PVDF membrane. Binding of the  $^{45}\text{Ca}^{2+}$  to the pc43 fusion protein was detected by autoradiography and was determined to be nearly as efficient as binding of  $^{45}\text{Ca}^{2+}$  to calmodulin. In contrast, there was no binding of calcium to purified

maltose binding protein lacking the pc43 extracellular domain. The pc43 subdomains EC-3 through EC-5 contain sequences highly homologous to the putative  $\text{Ca}^{2+}$  binding motifs found in E-cadherin. [See, Ringwald *et al.*, *EMBO J.*, 6: 3647-3653 (1987).]

5

#### Example 10

The expression of mRNA encoding pc42 and pc43 was assayed in various tissues and cell lines by Northern blot.

10

Total RNAs were prepared by the guanidium isothiocyanate method and poly(A)+ RNAs were isolated using a FastTrack kit (Invitrogen). RNA preparations were electrophoresed in a 0.8% agarose gel under denaturing conditions and transferred onto a nitrocellulose filter using a capillary method. Northern blot analyses were performed according to the method of Thomas, *Proc. Natl. Acad. Sci. USA*, 77: 5201-5205 (1980). The final wash was in 0.2X standard saline citrate containing 0.1% sodium dodecyl sulfate at 65°C for 10 minutes.

15

#### Protocadherin mRNA Expression in Adult Rat Tissues

20

Total mRNA preparations of rat tissues including brain, heart, liver, lung, skin, kidney and muscle were separated electrophoretically under denaturing conditions (10  $\mu\text{g}$  mRNA/lane) and transferred onto nitrocellulose filters. The filters were hybridized with  $^{32}\text{P}$ -labelled cDNA fragments MOUSE-326 (which corresponds to EC-4 of human pc42) and RAT-218 (which corresponds to EC-5 of human pc43). The mRNAs of both protocadherins were highly expressed in brain. The pc42 probe detected a major band of 7 kb and a minor band of 4 kb in size, possibly representing the products of alternative splicing. The pc43 probe hybridized to a major band of 5 kb in size and with minor bands of smaller sizes.

25

#### Developmental Expression of Protocadherin mRNA in Rat Brain

To examine the developmental regulation of mRNA expression of the protocadherins, brain mRNA from rats at embryonic days 17 and 20, neonatal

days 5 and 11 and from adult rats was prepared and subjected to Northern blot analysis as described above for other rat tissues.  $\beta$ -actin was used as an internal standard. mRNA levels for pc42 and pc43 proteins increased during embryonic development of the brain as compared with  $\beta$ -actin expression.

5 Protocadherin mRNA Expression in Human Cell Lines

Several neuronal and glial cell lines (including human SK-N-SH neuroblastoma, human U251 glioma, and mouse Neuro-2a neuroblastoma cell lines) were assayed by Northern blot using  $^{32}$ P-labelled for expression of pc42 and pc43 mRNA. Human cell lines were probed with HUMAN-42 (which corresponds to EC-4 of human pc42) and HUMAN-43 (which corresponds to EC-5 of human pc43) cDNA fragments while the mouse cell line was probed with MOUSE-326 (which corresponds to EC-4 of human pc42) and RAT-322 (which corresponds to EC-5 of human pc43) cDNA fragments. SK-N-SH human neuroblastoma cells and U251 human glioma cells were found to express pc43 mRNA and Neuro-2a mouse neuroblastoma cells were found to express pc42 mRNA.

Example 11

Expression of pc43 protein in various tissues, extracts and cells was assayed by Western blot and immunofluorescence microscopy.

20 Expression in Rat Cardiac Muscle Extracts

A rat heart non-ionic detergent extract was prepared by freezing a heart in liquid nitrogen after removal, powdering in a mortar and pestle, grinding briefly in a polytron in 0.5% Nonidet P40 in [10 mM PIPES (pH 6.8), 50 mM NaCl, 250 mM  $\text{NH}_4\text{SO}_4$ , 300 mM sucrose, 3 mM  $\text{MgCl}_2$ ] and microfuging for 15 minutes. Samples were separated by SDS/PAGE and electrophoretically transferred to nitrocellulose (Towbin *et al.*, *PNAS* 76:4350-4354, 1979). Two pc43 protein bands with molecular weights of 150 KDa and 140 KDa were

detected with rabbit polyclonal antibodies to pc43 by the immunoblot method described in Example 7.

#### Expression in Tissue Sections and Cells

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5 To determine the localization of the protocadherins in various tissues, human and rat adult tissues were removed, incubated in 30% sucrose in PBS for 30 minutes at 4°C, embedded in OCT compound (Tissue-Tek, Elkhart, Indiana) in cryomolds and quickly frozen. Six micron sections were cut and placed on glass slides. The slides were washed with PBS and fixed in 3% p-formaldehyde for 5 minutes. To permeabilize the tissue sections, the slides were  
10 immersed in -20°C acetone for 10 minutes and air dried. The sections were blocked with 2% goat serum and 1% BSA in PBS for 30 minutes and then incubated with the rabbit anti-pc43 polyclonal antisera for 1 hour at room temperature. The sections were rinsed 3 times in PBS containing 0.1% BSA and  
15 incubated with a biotinylated anti-rabbit (Vector Laboratories, Burlingame, California) in 1% BSA in PBS for 30 minutes. After rinsing 3 times, streptavidin-conjugated with FITC (Vector Laboratories) was added for 30 minutes and again washed 3 times. For co-localization studies, an appropriate primary antibody was used with a TRITC-conjugated secondary antibody.

#### A. Muscle

20 Immunolocalization of pc43 in rat cardiac muscle shows that pc43 is localized in a repeating pattern which is consistent with pc43 being associated with the sarcomeres. Sarcomeres are repetitive contractile units between the *fascia adherens* in skeletal and cardiac muscle. Co-localization with cytoskeletal proteins shows that pc43 is present at the ends of the sarcomeres in the Z lines  
25 which are associated with desmin and the actin-binding protein vinculin, and alpha-actinin. The thin microfilaments of F-actin are associated with the thick myosin filaments between the Z lines. In contrast, N-cadherin is localized at the ends of cardiac myocytes at the *fascia adherens* junctions at sites of myocyte:myocyte contact. The localization of pc43 in cardiac muscle suggests

that pc43 may play a role in muscle contraction in the anchoring of the contractile apparatus to the plasma membrane.

5 Similar localization for pc43 was observed in rat skeletal muscle. Ultrastructural studies have shown that dystrophin, the gene product lacking in Duchenne muscular dystrophy, is a component of the sarcolemma [Porter *et al.*, *J. Cell. Biol.*, 117:997-1005 (1992)]. The sarcolemma is connected to the contractile apparatus at the M and Z lines where pc43 is localized.

#### B. Brain

10 Reactivity of anti-pc43 polyclonal antibody and monoclonal antibody 38I2C on frozen sections of rat and human cerebellum, respectively, shows that the major sites of pc43 expression are located in Purkinje cells and the granule cell layer which contains numerous small neurons.

#### C. Placenta

15 Strong reactivity of monoclonal antibody 38I2C with human syncytiotrophoblasts was also observed in development of the placenta at an early state (5-7 weeks of gestation). Expression appeared to gradually decrease as the stage progressed indicating that pc43 may be involved in the implantation of fertilized eggs into the placenta.

#### D. Neuroblastoma and Astrocytoma Cells

20 Immunocytochemical localization of pc43 in Sk-N-SH neuroblastoma cells and UW28 astrocytoma cells using anti-pc43 antibodies reveals a punctate cell surface distribution of pc43 and in some cells there is a localization at the tips of extensions of neuronal foot processes. At sites of cell-cell contact of UW28 astrocytoma cells, pc43 is organized in a series of parallel  
25 lines. The lines start at the contact site and extend approximately 5 micron. F-actin microfilaments were identified with rhodamine-phalloidin (Molecular Probes, Eugene, Oregon, as described by the manufacturer) showing that the microfilaments in the cell appear to end in the pc43 linear structures which extend from the edge of the cell at sites of cell contact.

Immunoblotting studies with pc43 specific antibodies show that a protein with a molecular weight of 140 kDa is recognized in human Sk-N-SH neuroblastoma cells and in UW28 astrocytoma cells.

#### E. Osteoblasts

5 Immunocytochemical localization of pc43 using monoclonal antibody 38I2C in tow human ostogenic sarcoma cell lines [SaOS (ATCC HTB 85) and MG-63 (ATCC CRL 1427)] and in cultures of normal human trabecular osteoblasts [culture system described in Civitelli *et al.*, *J. Clin. Invest.*, 91: 1888-1896 (1993)] showed that pc43 is expressed in osteoblasts in a pattern similar to  
10 that seen in UW28 astrocytoma cells. At sites of cell-cell contact, pc43 is organized in a series of parallel lines that appear to correspond to the actin stress fibers. In addition, in some cells, pc43 appears to localize at the tips of contacting cell processes. Northern blot analysis provides additional evidence that pc43 is expressed in normal human trabecular osteoblasts. A pc43 specific DNA  
15 probe hybridized to a major band of 5 kb in samples of poly-A mRNA isolated from normal human trabecular osteoblasts.

#### Example 12

*In situ* hybridization experiments using protocadherin specific RNA probes were performed on cryosections of rat tissue.

20 Sense and antisense <sup>35</sup>S-riboprobes were made using the standard procedure described by Promega (Madison, Wisconsin). An approximately 400 bp EcoRI-XbaI fragment of the MOUSE-326 cDNA clone was used as a pc42 specific probe. This fragment encodes the middle of EC-3 to the end of EC-4 of pc42. An approximately 700 bp SmaI fragment of the RAT-218 cDNA clone was  
25 used as a pc43 specific probe. The fragment encodes the end of EC-3 to the end of EC-5 of pc43.

Rat adult tissues were harvested and immediately embedded with OCT Compound (Tissue-Tek) in cryomolds and quickly frozen in a bath of 95% ethanol/dry ice. The frozen blocks were stored at -80°C until cut. Six micron

tissue sections were cut using a cryostat (Reichert-Jung, Model #2800 Frigocut N, Leica, Inc., Gilroy, California). Cut tissue sections were stored at -80°C.

5 The *in situ* protocol used was a variation of that described by Angerer *et al.*, *Methods in Enzymology*, 152: 649-660, (1987). All solutions were treated with diethylpyrocarbonate (DEPC, Sigma, St. Louis, Missouri) to remove RNase contamination. The tissue sections were first fixed in 4% paraformaldehyde at 4°C for 20 minutes. To remove excess paraformaldehyde and stop the tissue fixation, the slides were washed in PBS (phosphate buffered saline), denatured in a graded series of alcohols (70, 95, 100%) and then dried. 10 To prevent the tissue from detaching from the glass slide during the *in situ* procedure, the tissue sections were treated in a poly-L-lysine solution (Sigma) at room temperature for 10 minutes. To denature all RNA in the tissue, the sections were placed in a solution of 70% formamide/2x SSC (0.15 M NaCl/0.3 M Na citrate, pH 7.0) at 70°C for 2 minutes after which they were rinsed in chilled 2x 15 SSC, dehydrated in a graded series of alcohols and then dried. Once dried, the sections were prehybridized in hybridization buffer [50% formamide/50 mM DTT (dithiothrietol)/0.3M NaCl/20 mM Tris, pH 8.0/5 mM EDTA/1X Denhardt's (0.02% Ficoll Type 400/0.02% polyvinylpyrrolidone/0.02% BSA)/10% Dextran Sulfate] at the final hybridization temperature for approximately 4 hours. After 20 prehybridization, approximately  $1 \times 10^6$  cpm of the appropriate riboprobe was added to each section. The sections were generally hybridized at 45°C overnight (12-16 hours). To insure that the hybridization seen was specific, in some experiments the hybridization stringency was increased by raising the hybridization temperature to 50°C. As both the 45°C and 50°C experiments gave 25 comparable results, the standard hybridization temperature used was 45°C.

To remove excess, nonhybridized probe, the sections were put through a series of washes. The sections were first rinsed in 4X SSC to remove the bulk of the hybridization solution and probe. Next a 15 minute wash in 4X SSC/50 mM DTT was carried out at room temperature. Washes at increased

stringencies were also utilized. A 40 minute wash in 50% formamide/2X SSC/50 mM DTT was performed at 60°C. Four final room temperature washes were carried out for 10 minutes each: two in 2X SSC and two in 0.1X SSC. The washed slides were dehydrated in a graded series of alcohols and dried.

5 To visualize the hybridized probe, the slides were dipped in Kodak NTB2 nuclear emulsion (International Biotechnology, New Haven, Connecticut) which had been diluted 1:1 in dH<sub>2</sub>O. Once dry, the slides were stored at 4°C in light-tight boxes for the appropriate exposure time. The *in situ* slides were independently viewed by two persons and scored positive or negative for  
10 hybridization signal.

All *in situ* hybridization studies were performed on rat tissue. Because results from Northern blot experiments (see Example 9) indicated that both pc42 and pc43 are expressed in adult brain, *in situ* hybridization studies were carried out to localize the expression of these molecules to specific brain cell  
15 types. Hybridization seen in the normal adult rat brain was specific (no background hybridization was seen with the sense probes) and was localized to specific regions in the brain. The overall pattern of expression seen for pc42 and pc43 was very similar, with the major difference being in the level of expression. pc43 appears to be expressed at a lower level than pc42. Both molecules are  
20 expressed in the germinal and pyramidal cells of the hippocampus, Purkinje cells of the cerebellum and neurons in grey matter. In addition, pc42 is expressed in glial cells in the white matter but, in contrast to the expression of pc43 in glioma cell lines (as described in Example 9), expression of pc43 in normal glial cells was not observed. In the spinal chord, both protocadherins are expressed in the  
25 motor neurons in the gray matter and pc42 is expressed in the glial cells in the white matter.

When expression of both protocadherin molecules was analyzed in brains and spinal chords from rats having EAE (experimental allergic encephalomyelitis) [Vandenbark et al., *Cell. Immunol.*, 12: 85-93 (1974)], the  
30 same structures as described above were found to be positive. In addition,



expression of pc42 was observed in the leukocytic infiltrates in the EAE tissues. Expression of pc42 in leukocytes was confirmed by *in situ* hybridization analysis of two leukocytic cell lines, RBL-1 and y3.

Expression of both protocadherin-42 and -43 was observed in the developing brain of rat embryos at all embryological days tested (E15-E19). In addition protocadherin-43 was observed in the developing rat heart at all embryological days tested (E13-E19). This finding is consistent with the immunohistochemistry results showing protocadherin-43 expression in adult heart.

To determine possible roles of protocadherins in the development of the nervous system, expression profiles of protocadherin members in developing rat brain and adult rat brain were also examined by *in situ* hybridization. A series of coronal, sagittal and horizontal sections of rat brains at postnatal days 0, 6, 14, 30 (P0 through P30) and at 3 months (young adult) were hybridized with labelled cRNA probes corresponding to various protocadherins of the invention including pc42, pc43, RAT-212, RAT-411, and RAT-418. In developing brain, RAT-411 was expressed at high levels in neurons of the olfactory bulb, *i.e.*, mitral cells and periglomerular cells. The expression of RAT-411 mRNA was transient; expression appeared at P0, peaked at P6, diminished by P14, and was undetectable at P30 and in adult brain. In the adult, pc43 mRNA was found to be expressed predominantly in Purkinje cells in the cerebellum. The expression of pc43 mRNA in Purkinje cells was observed from the beginning of Purkinje cell differentiation at around P6. Other protocadherin members were expressed at very low levels in various areas of developing and adult brains. These results indicate that protocadherin members are differentially expressed during the development of the central nervous system, and suggest that RAT-411 and pc43 have specific roles during the development of olfactory bulb neurons and Purkinje cells, respectively.

**Example 13**

Conventional immunoprecipitations using pc43-specific polyclonal antibodies and monoclonal antibody 38I2C were performed to identify proteins that interacted with pc43 in L cell transfectants.

5           The pc43 and chimeric pc43 transfectants were metabolically labeled by incubating the cells in Dulbecco's modified Eagle's medium containing [35S] methionine (50 uCi/ml) overnight. After washing, the transfectants were lysed with PBS containing Triton X 100 and incubated with anti-pc43 antibody. The immunocomplexes were then collected using protein A-Sepharose beads. The  
10       resulting beads were washed five times with a washing buffer (50mM Tris-HCl, pH 8.0, containing 0.5M NaCl, 0.1% ovalbumin, 0.5% NP-40, 0.5% Triton X 100 and 1mM EDTA) at room temperature. Protein was separated by SDS-PAGE and subjected to autoradiography.

15           The chimeric pc43 co-precipitated with 105 kDa and a 95 kDa bands that are likely to correspond to  $\alpha$ - and  $\beta$ -catenins, respectively, because anti- $\alpha$ -catenin and anti- $\beta$ -catenin antibodies stained comparable bands. Pc43, on the other hand, co-precipitated with a 120 kDa band.

20           While the present invention has been described in terms of specific methods and compositions, it is understood that variations and modifications will occur to those skilled in the art. Therefore, only such limitations as appear in the claims should be placed on the invention.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT: Suzuki, Shintaro
- (ii) TITLE OF INVENTION: Protocadherin Materials and Methods
- (iii) NUMBER OF SEQUENCES: 115
- (iv) CORRESPONDENCE ADDRESS:  
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- (v) COMPUTER READABLE FORM:  
    (A) MEDIUM TYPE: Floppy disk  
    (B) COMPUTER: IBM PC compatible  
    (C) OPERATING SYSTEM: PC-DOS/MS-DOS  
    (D) SOFTWARE: PatentIn Release #1.0, Version #1.25
- (vi) CURRENT APPLICATION DATA:  
    (A) APPLICATION NUMBER:  
    (B) FILING DATE:  
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- (vii) PRIOR APPLICATION DATA  
    (A) APPLICATION NUMBER: PCT/US93/12588  
    (B) FILING DATE: 23 DEC 1993
- (vii) PRIOR APPLICATION DATA  
    (A) APPLICATION NUMBER: US 07/998,003  
    (B) FILING DATE: 29 DEC 1992
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(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 17 base pairs  
    (B) TYPE: nucleic acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA

09860573 064301

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

AARSSNNTNG AYTRYGA

17

(2) INFORMATION FOR SEQ ID NO:2:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 17 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

TTRCTRTTRC GNGGNNN

17

(2) INFORMATION FOR SEQ ID NO:3:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 131 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

AAGGGAGTGG ACTTTGAGGA GCAGCCTGAG CTTAGTCTCA TCCTCACGGC TTTGGATGGA	60
GGGACTCCAT CCAGGTCTGG GACTGCATTG GTTCAAGTGG AAGTCATAGA TGCCAATGAC	120
AACGCACCGT A	131

(2) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

Lys	Gly	Val	Asp	Phe	Glu	Glu	Gln	Pro	Glu	Leu	Ser	Leu	Ile	Leu	Thr
1				5					10					15	
Ala	Leu	Asp	Gly	Gly	Thr	Pro	Ser	Arg	Ser	Gly	Thr	Ala	Leu	Val	Gln
		20						25					30		
Val	Glu	Val	Ile	Asp	Ala	Asn	Asp	Asn	Ala	Pro					
		35						40							

(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

AAACGCATGG ATTCGAGGA GTCTTCCTCC TACCAGATCT ATGTGCAAGC TACTGACCGG	60
GGACCACTAC CCATGGCGGG TCATTGCAAG GTGTTGGTGG ACATTATAGA TGTGAACGAC	120
AACGCACCTA A	131

(2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 43 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

Lys	Ala	Met	Asp	Phe	Glu	Glu	Ser	Ser	Ser	Tyr	Gln	Ile	Tyr	Val	Gln
1				5					10					15	
Ala	Thr	Asp	Arg	Gly	Pro	Val	Pro	Met	Ala	Gly	His	Cys	Lys	Val	Leu
			20					25					30		
Val	Asp	Ile	Ile	Asp	Val	Asn	Asp	Asn	Ala	Pro					
		35					40								

(2) INFORMATION FOR SEQ ID NO:7:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

AAGCGACTGG ACTTTGAGAC CCTGCAGACC TTCGAGTTCA GCGTGGGTGC CACAGACCAT	60
GGCTCCCCCT CGCTCCGCAG TCAGGCTCTG GTGCGCGTGG TGGTGCTGGA CCACAATGAC	120

09860573.064301

AATGCCCCCA A

131

(2) INFORMATION FOR SEQ ID NO:8:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

Lys	Arg	Leu	Asp	Phe	Glu	Thr	Leu	Gln	Thr	Phe	Glu	Phe	Ser	Val	Gly
1				5				10						15	
Ala	Thr	Asp	His	Gly	Ser	Pro	Ser	Leu	Arg	Ser	Gln	Ala	Leu	Val	Arg
			20					25					30		
Val	Val	Val	Leu	Asp	His	Asn	Asp	Asn	Ala	Pro					
		35				40									

(2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 131 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

AAGGGCCTGG	ATTACGAGGC	ACTGCAGTCC	TTCGAGTTCT	ACGTGGGCGC	TACAGATGGA	60
GGCTCACCCG	CGCTCAGCAG	CCAGACTCTG	GTGCGGATGG	TGGTGCTGGA	TGACAACGAC	120
AACGCCCCCTA	A					131

(2) INFORMATION FOR SEQ ID NO:10:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:-

Lys	Gly	Leu	Asp	Tyr	Glu	Ala	Leu	Gln	Ser	Phe	Glu	Phe	Tyr	Val	Gly
1				5				10					15		

Met Val Val Leu Asp Asp Asn Asp Asn Ala Pro  
35 40

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 131 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

AAGGCGTTTG	ATTTTGAGGA	TCAGAGAGAG	TTCCAGCTAA	CCGCTCATAT	AAACGACGGA	60
GGTACCCCGG	TTTTGGCCAC	CAACATCAGC	GTGAACATAT	TTGTTACTGA	CCGCAATGAC	120
AACGCCCCGC	A					131

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 43 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

Lys Ala Phe Asp Phe Glu Asp Gln Arg Glu Phe Gln Leu Thr Ala His  
1 5 10 15  
Ile Asn Asp Gly Gly Thr Pro Val Leu Ala Thr Asn Ile Ser Val Asn  
20 25 30  
Ile Phe Val Thr Asp Arg Asn Asp Asn Ala Pro  
35 40

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

AAGGCGGTGG ATTACGAAAT CACCAAGTCC TATGAGATAG ATGTTCAAGC CCAAGATCTG 60  
GGTCCCAATT CTATTCCTGC TCATTGCAAA ATTATAATTA AGGTCGTGGA TGTCAACGAC 120  
AACGCTCCCA A 131

(2) INFORMATION FOR SEQ ID NO:14:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 43 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

Lys Ala Val Asp Tyr Glu Ile Thr Lys Ser Tyr Glu Ile Asp Val Gln  
1 5 10 15  
Ala Gln Asp Leu Gly Pro Asn Ser Ile Pro Ala His Cys Lys Ile Ile  
20 25 30  
Ile Lys Val Val Asp Val Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:15:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 135 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

TATGACCATG ATTACGAGAC AACCAAAGAA TATACACTGC GGATCCGGGC CCAGGATGGT 60  
GGCCGGACTC CACTTTCCAA CGTCTCCGGT CTAGTAACCG TGCAGGTCCT AGACATCAAC 120  
GACAATGCCC CCCC 135

(2) INFORMATION FOR SEQ ID NO:16:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 44 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein



(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

Tyr Asp His Asp Tyr Glu Thr Thr Lys Glu Tyr Thr Leu Arg Ile Arg  
 1 5 10 15  
 Ala Gln Asp Gly Gly Arg Thr Pro Leu Ser Asn Val Ser Gly Leu Val  
 20 25 30  
 Thr Val Gln Val Leu Asp Ile Asn Asp Asn Ala Pro  
 35 40

(2) INFORMATION FOR SEQ ID NO:17:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 129 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

GGGGGGTCGA TTACGAGGAG AACGGCATGT TAGAGATCGA CGTGCAGGCC AGAGACCTAG 60  
 GACCTAACCC AATTCCAGCC CATTGCAAGG TCACAGTCAA GCTCATCGAC CGCAATGATA 120  
 ACGCCCCCA 129

(2) INFORMATION FOR SEQ ID NO:18:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 43 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

Arg Gly Val Asp Tyr Glu Glu Asn Gly Met Leu Glu Ile Asp Val Gln  
 1 5 10 15  
 Ala Arg Asp Leu Gly Pro Asn Pro Ile Pro Ala His Cys Lys Val Thr  
 20 25 30  
 Val Lys Leu Ile Asp Arg Asn Asp Asn Ala Pro  
 35 40

0988057.061304

(2) INFORMATION FOR SEQ ID NO:19:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 131 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

AAGGGGTTGG ACTACGAAGA CACCAAATC CATGAGATTT ACATCCAGGC CAAAGACAAA	60
GGTGCCAATC CGGAAGGAGC GCATTGCAAA GTACTGGTAG AGGTTGTGGA CGTTAACGAC	120
AATGCCCTC A	131

(2) INFORMATION FOR SEQ ID NO:20:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

Lys Gly Leu Asp Tyr Glu Asp Thr Lys Leu His Glu Ile Tyr Ile Gln	
1 5 10 15	
Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys Val Leu	
20 25 30	
Val Glu Val Val Asp Val Asn Asp Asn Ala Pro	
35 40	

(2) INFORMATION FOR SEQ ID NO:21:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 131 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

AAGGGTTTGG ACTTTGAGCA AGTAGATGTC TACAAAATCC GCGTTGACGC GACGGACAAA	60
GGACACCTC CGATGGCAGG CCATTGCACT GTTTTAGTGA GGGTATTGGA TGAAAACGAC	120

09880573 061301

AATGCGCCTC T

131

(2) INFORMATION FOR SEQ ID NO:22:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

Lys	Gly	Leu	Asp	Phe	Glu	Gln	Val	Asp	Val	Tyr	Lys	Ile	Arg	Val	Asp	
1					5				10					15		
Ala	Thr	Asp	Lys	Gly	His	Pro	Pro	Met	Ala	Gly	His	Cys	Thr	Val	Leu	
			20					25					30			
Val	Arg	Val	Leu	Asp	Glu	Asn	Asp	Asn	Ala	Pro						
			35					40								

(2) INFORMATION FOR SEQ ID NO:23:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 134 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

AAGGGTATAG	ACTTCGAGCA	GATCAAGGAC	TTCAGCTTTC	AAGTGAAGC	CCGGGACGCC	60
GGCAGTCCCC	AGGCGCTGTC	CGGCAACTGC	ACTGTCAACA	TCTTGATAGT	GGATCAGAAC	120
GACAACGCCC	CTAA					134

(2) INFORMATION FOR SEQ ID NO:24:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 44 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

Lys	Gly	Ile	Asp	Phe	Glu	Gln	Ile	Lys	Asp	Phe	Ser	Phe	Gln	Val	Glu
1				5				10					15		

- 44 -

Ala Arg Asp Ala Gly Ser Pro Gln Ala Leu Ala Gly Asn Thr Thr Val  
20 25 30

Asn Ile Leu Ile Val Asp Gln Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:25:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 134 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

AAGCCGTTTCG ACTATGAGCA AACCGCCAAC ACGCTGGCAC AGATTGACGC CGTGCTGGAA 60  
AAACAGGGCA GCAATAAATC GAGCATTCTG GATGCCACCA TTTTCCTGGC CGATAAAAAC 120  
GACAATGCGC CAGA 134

(2) INFORMATION FOR SEQ ID NO:26:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 44 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

Lys Pro Phe Asp Tyr Glu Gln Thr Ala Asn Thr Leu Ala Gln Ile Asp  
1 5 10 15  
Ala Val Leu Glu Lys Gln Gly Ser Asn Lys Ser Ser Ile Leu Asp Ala  
20 25 30  
Thr Ile Phe Leu Ala Asp Lys Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:27:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 131 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

AAGCGGCTGG ATTTGGAACA GTTCCAGCAG CACAAGCTGC TCGTAAGGGC TGTGATGGA 60  
GGAATGCCGC CACTGAGCAG CGATGTGGTC GTCAGTGTGG ATGTCACCGA CCTCAACGAT 120  
AACGCGCCCT A 131

(2) INFORMATION FOR SEQ ID NO:28:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 43 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

Lys Arg Leu Asp Phe Glu Gln Phe Gln Gln His Lys Leu Leu Val Arg  
1 5 10 15  
Ala Val Asp Gly Gly Met Pro Pro Leu Ser Ser Asp Val Val Val Thr  
20 25 30  
Val Asp Val Thr Asp Leu Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:29:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 131 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

AAGGGGATAG ACTTTGAGAG TGAGAATTAC TATGAATTTG ATGTGCGGGC TCGCGATGGG 60  
GGTTCTCCAG CCATGGAGCA ACATTGCAGC CTTGAGTGG ATCTGCTGGA CGTAAATGAC 120  
AACGCCCCAC T 131

(2) INFORMATION FOR SEQ ID NO:30:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 43 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

09680573-061301

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

Lys Gly Ile Asp Phe Glu Ser Glu Asn Tyr Tyr Glu Phe Asp Val Arg  
 1 5 10 15  
 Ala Arg Asp Gly Gly Ser Pro Ala Met Glu Gln His Cys Ser Leu Arg  
 20 25 30  
 Val Asp Leu Leu Asp Val Asn Asp Asn Ala Pro  
 35 40

(2) INFORMATION FOR SEQ ID NO:31:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

AAGGCATTGG ACTTTGAGGC CCGGCGACTG TATTCGCTGA CAGTTCAGGC CACGGACCGA 60  
 GCGGTGCCCT CGCTACCGG GCGTGCCGAA GCGCTTATCC AGCTGCTAGA TGTCAACGAC 120  
 AACGCACCCA T 131

(2) INFORMATION FOR SEQ ID NO:32:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 43 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:

Lys Ala Leu Asp Phe Glu Ala Arg Arg Leu Tyr Ser Leu Thr Val Gln  
 1 5 10 15  
 Ala Thr Asp Arg Gly Val Pro Ser Leu Thr Gly Arg Ala Glu Ala Leu  
 20 25 30  
 Ile Gln Leu Leu Asp Val Asn Asp Asn Ala Pro  
 35 40

(2) INFORMATION FOR SEQ ID NO:33:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 125 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

AAGCCAATTG ATTACGAGGC AACTCCATAC TATAACATGG AAATTGTAGC CACAGACAGC	60
GGAGGTCTTT CGGGAAAATG CACTGTGTCT ATACAGGTGG TGGATGTGAA CGACAACGCC	120
CCCAA	125

(2) INFORMATION FOR SEQ ID NO:34:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 41 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:

Lys	Pro	Ile	Asp	Tyr	Glu	Ala	Thr	Pro	Tyr	Tyr	Asn	Met	Glu	Ile	Val
1				5					10					15	
Ala	Thr	Asp	Ser	Gly	Gly	Leu	Ser	Gly	Lys	Cys	Thr	Val	Ser	Ile	Gln
			20					25					30		
Val	Val	Asp	Val	Asn	Asp	Asn	Ala	Pro							
		35					40								

(2) INFORMATION FOR SEQ ID NO:35:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 446 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

AAGCGGGTAG ACTTCGAAAT GTGCAAAGA TTTTACCTTG TGGTGGAAGC TAAAGACGGA	60
GGCACCCCAG CCCTCAGCAC GGCAGCCACT GTCAGCATCG ACCTCACAGA TGTGAATGAT	120

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AACCCTCCTC GGTTCAGCCA AGATGTCTAC AGTGCTGTCA TCAGTGAGGA TGCCTTAGAG 180  
 GGGGACTCTG TCATTCTGCT GATAGCAGAA GATGTGGATA GCAAGCCTAA TGGACAGATT 240  
 CGGTTTTCCA TCGTGGGTGG AGATAGGGAC AATGAATTTG CTGTGATCC AATCTTGGGA 300  
 CTTGTGAAAG TTAAGAAGAA ACTGGACCGG GAGCGGGTGT CAGGATACTC CCTGCTCATC 360  
 CAGGCAGTAG ATAGTGGCAT TCCTGCAATG TCCTCAACGA CAACTGTCAA CATTGATATT 420  
 TCTGATGTGA ACGACAACGC CCCCCT 446

(2) INFORMATION FOR SEQ ID NO:36:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 148 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

Lys Arg Val Asp Phe Glu Met Cys Lys Arg Phe Tyr Leu Val Val Glu  
 1 5 10 15  
 Ala Lys Asp Gly Gly Thr Pro Ala Leu Ser Thr Ala Ala Thr Val Ser  
 20 25 30  
 Ile Asp Leu Thr Asp Val Asn Asp Asn Pro Pro Arg Phe Ser Gln Asp  
 35 40 45  
 Val Tyr Asp Ala Val Ile Ser Glu Asp Ala Leu Glu Gly Asp Ser Val  
 50 55 60  
 Ile Leu Leu Ile Ala Glu Asp Val Asp Ser Lys Pro Asn Gly Gln Ile  
 65 70 75 80  
 Arg Phe Ser Ile Val Gly Gly Asp Arg Asp Asn Glu Phe Ala Val Asp  
 85 90 95  
 Pro Ile Leu Gly Leu Val Lys Val Lys Lys Lys Leu Asp Arg Glu Arg  
 100 105 110  
 Val Ser Gly Tyr Ser Leu Leu Ile Gln Ala Val Asp Ser Gly Ile Pro  
 115 120 125  
 Ala Met Ser Ser Thr Thr Thr Val Asn Ile Asp Ile Ser Asp Val Asn  
 130 135 140  
 Asp Asn Ala Pro  
 145

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(2) INFORMATION FOR SEQ ID NO:37:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 440 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

AAGGGGGTTG ATTATGAGAC AAACCCACGG CTACGACTGG TGCTACAGGC AGAGAGTGGA 60  
 GGAGCCTTTG CTTTCTCGGT GCTGACCCTG ACCCTTCAAG ATGCCAATGA CAATGCTCCC 120  
 CGTTTCCTGC AGCCTCACTA CGTGGCTTTC CTGCCAGAGT CCCGACCCTT GGAAGGGCCC 180  
 CTGCTGCAGG TGAAGCAGA CGACCTGGAT CAAGGCTCTG GAGGACAGAT CTCCTACAGT 240  
 CTGGCTGCAT CCCAGCCAGC ACGGGGCTTG TTCCATGTAG ACCCAGCCAC AGGCACTATC 300  
 ACTACCACAG CCATCCTGGA CCGGAAATC TGGGCTGAAA CACGGCTGGT ACTGATGGCC 360  
 ACAGACAGAG GAAGCCCAGC ATTGCTGGGC TCAGCTACCC TGACAGTGAT GGTCATCGAT 420  
 ACCAACGACA ATGCTCCCCT 440

(2) INFORMATION FOR SEQ ID NO:38:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 146 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:

Lys Gly Val Asp Tyr Glu Thr Asn Pro Arg Leu Arg Leu Val Leu Gln  
 1 5 10 15  
 Ala Glu Ser Gly Gly Ala Phe Ala Phe Ser Val Leu Thr Leu Thr Leu  
 20 25 30  
 Gln Asp Ala Asn Asp Asn Ala Pro Arg Phe Leu Gln Pro His Tyr Val  
 35 40 45  
 Ala Phe Leu Pro Glu Ser Arg Pro Leu Glu Gly Pro Leu Leu Gln Val  
 50 55 60  
 Glu Ala Asn Asp Leu Asp Gln Gly Ser Gly Gly Gln Ile Ser Tyr Ser  
 65 70 75 80  
 Leu Ala Ala Ser Gln Pro Ala Arg Gly Leu Phe His Val Asp Pro Ala  
 85 90 95

Thr Gly Thr Ile Thr Thr Thr Ala Ile Leu Asp Arg Glu Ile Trp Ala  
 100 105 110  
 Glu Thr Arg Leu Val Leu Met Ala Thr Asp Arg Gly Ser Pro Ala Leu  
 115 120 125  
 Val Gly Ser Ala Thr Leu Thr Val Met Val Ile Asp Thr Asn Asp Asn  
 130 135 140  
 Ala Pro  
 145

(2) INFORMATION FOR SEQ ID NO:39:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 124 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

AAGGTCTCGA TTATGAGGCA ACTCCATATT ATAACGTGGA AATTGTAGCC ACAGATGGTG 60  
 GGGGCCTTTC AGGAAAATGC ACTGTGGCTA TAGAAGTGGT GGATGTGAAC GACGGCGCTC 120  
 CAAT 124

(2) INFORMATION FOR SEQ ID NO:40:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 41 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:40:

Lys Gly Leu Asp Tyr Glu Ala Thr Pro Tyr Tyr Asn Val Glu Ile Val  
 1 5 10 15  
 Ala Thr Asp Gly Gly Ala Phe Asp Glu Asn Cys Thr Val Ala Ile Glu  
 20 25 30  
 Val Val Asp Val Asn Asp Asn Ala Pro  
 35 40

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(2) INFORMATION FOR SEQ ID NO:41:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 8 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:41:

Asp Xaa Asn Glu Xaa Pro Xaa Phe  
1 5

(2) INFORMATION FOR SEQ ID NO:42:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 8 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:42:

Asp Xaa Asp Glu Xaa Pro Xaa Phe  
1 5

(2) INFORMATION FOR SEQ ID NO:43:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 9 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:43:

Asp Xaa Asn Asp Asn Xaa Pro Xaa Phe  
1 5

(2) INFORMATION FOR SEQ ID NO:44:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 131 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:44:

AAGCGGATGG ATTTTGAAGA CACCAAACCTC CATGAGATTT ACATCCAGGC CAAAGACAAA 60  
GGTGCCAATC CCGAAGGAGC GCATTGCAAA GTACTTGTAG AGGTTGTAGA CGTAAACGAC 120  
AACGCCCCAG T 131

(2) INFORMATION FOR SEQ ID NO:45:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 43 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:45:

Leu Arg Met Asp Phe Glu Asp Thr Lys Leu His Glu Ile Tyr Ile Gln  
1 5 10 15  
Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys Val Leu  
20 25 30  
Val Glu Val Val Asp Val Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:46:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 131 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:46:

AAGGCTTTGG ATTACGAGGA TCAGAGAGAG TTCCAACATA CAGCTCATAT AAACGACGGA 60  
GGTACCCCAG TCTTAGCCAC CAACATCAGC GTGAACGTAT TTGTTACTGA CCGCAATGAT 120  
AACGCCCCCT A 131

(2) INFORMATION FOR SEQ ID NO:47:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 43 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:47:

Lys Ala Leu Asp Tyr Glu Asp Gln Arg Glu Phe Gln Leu Thr Ala His  
 1 5 10 15  
 Ile Asn Asp Gly Gly Thr Pro Val Leu Ala Thr Asn Ile Ser Val Asn  
 20 25 30  
 Val Phe Val Thr Asp Arg Asn Asp Asn Ala Pro  
 35 40

(2) INFORMATION FOR SEQ ID NO:48:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:48:

AAGCGCTTGG ACTACGAGGA GAGTAACAAT TATGAAATTC ACGTGGATGC TACAGATAAA 60  
 GGATACCCAC CTATGGTTGC TCACTGCACC GTACTCGTGG GAATCTTGGA TGAAAATGAC 120  
 AACGCACCCA T 131

(2) INFORMATION FOR SEQ ID NO:49:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 43 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:49:

Lys Arg Leu Asp Tyr Glu Glu Ser Asn Asn Tyr Glu Ile His Val Asp  
 1 5 10 15  
 Ala Thr Asp Lys Gly Tyr Pro Pro Met Val Ala His Cys Thr Val Leu  
 20 25 30  
 Val Gly Ile Leu Asp Glu Asn Asp Asn Ala Pro  
 35 40

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(2) INFORMATION FOR SEQ ID NO:50:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 131 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:50:

AAACCGGTGG ACTACGAGAA AGTCAAAGAC TATACCATCG AGATCGTGGC TGTGGATTCC	60
GGCAACCCTC CACTCTCTAG CACCAACTCC CTCAAGGTGC AGGTGGTAGA CGTCAACGAT	120
AACGCCCCCTC T	131

(2) INFORMATION FOR SEQ ID NO:51:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:51:

Lys	Pro	Val	Asp	Tyr	Glu	Lys	Val	Lys	Asp	Tyr	Thr	Ile	Glu	Ile	Val
1				5					10				15		
Ala	Val	Asp	Ser	Gly	Asn	Pro	Pro	Leu	Ser	Ser	Thr	Asn	Ser	Leu	Lys
			20					25				30			
Val	Gln	Val	Val	Asp	Val	Asn	Asp	Asn	Ala	Pro					
			35				40								

(2) INFORMATION FOR SEQ ID NO:52:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 131 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:52:

AAGCCTTTTG ATTTGAGGA CACCAAATC CATGAGATT ACATCCAGGC CAAAGACAAG	60
GGCGCCAATC CCGAAGGAGC ACATTGCAAA GTGTTGGTGG AGGTTGTGGA TGTGAACGAC	120

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AATGCCCCCTC A

131

(2) INFORMATION FOR SEQ ID NO:53:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:53:

Lys Pro Phe Asp Phe Glu Asp Thr Lys Leu His Glu Ile Tyr Ile Gln  
 1 5 10 15  
 Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys Val Leu  
 20 25 30  
 Val Glu Val Val Asp Val Asn Asp Asn Ala Pro  
 35 40

(2) INFORMATION FOR SEQ ID NO:54:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 122 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:54:

AAAGGTGTCG ATTACGAGGT GAGTCCACGG CTGCGACTGG TGCTGCAGGC AGAGAGTCGA 60  
 GGAGCCTTTG CCTTCACTGT GCTGACCCTG ACCCTGCAAG ATGCCAACGA CAACGCCCCG 120  
 AG 122

(2) INFORMATION FOR SEQ ID NO:55:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 40 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:55:

Lys Gly Val Asp Tyr Glu Val Ser Pro Arg Leu Arg Leu Val Leu Gln  
 1 5 10 15

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Ala Glu Ser Arg Gly Ala Phe Ala Phe Thr Val Leu Thr Leu Thr Leu  
20 25 30

Gln Asp Ala Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:56:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 131 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:56:

AAAGGGATTG ATTACGAGCA GTTGAGAGAC CTACAGCTGT GGGTGACAGC CAGCGACAGC 60  
GGGGACCCGC CTCTTAGCAG CAACGTGTCA CTGAGCCTGT TTGTGCTGGA CCAGAACGAC 120  
AACGCCCCCCC T 131

(2) INFORMATION FOR SEQ ID NO:57:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:57:

Lys Gly Ile Asp Tyr Glu Gln Leu Arg Asp Leu Gln Leu Trp Val Thr  
1 5 10 15  
Ala Ser Asp Ser Gly Asp Pro Pro Leu Ser Ser Asn Val Ser Leu Ser  
20 25 30  
Leu Phe Val Leu Asp Gln Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:58:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 125 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:58:

AAGGCGGTCG ATTTTGAGCG CACATCCTCT TATCAACTCA TCATTCAGGC CACCAATATG 60  
GCAGGAATGG CTTCCAATGC TACAGTCAAT ATTCAGATTG TTGATGAAAA CGACAACGCC 120  
CCCCA 125

(2) INFORMATION FOR SEQ ID NO:59:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 41 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:59:

Lys Ala Val Asp Phe Glu Arg Thr Ser Ser Tyr Gln Leu Ile Ile Gln  
1 5 10 15  
Ala Thr Asn Met Ala Gly Met Ala Ser Asn Ala Thr Val Asn Ile Gln  
20 25 30  
Ile Val Asp Glu Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:60:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 131 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:60:

AAACGGCTAG ACTTTGAAAA GATACAAAAA TATGTTGTAT GGATAGAGGC CAGAGATGGT 60  
GGTTTCCCTC CTTTCTCCTC TTACGAGAAA CTTGATATAA CAGTATTAGA TGTCAACGAT 120  
AACGCGCCTA A 131

(2) INFORMATION FOR SEQ ID NO:61:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 43 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:61:

Lys Arg Leu Asp Phe Glu Lys Ile Gln Lys Tyr Val Val Trp Ile Glu  
 1 5 10 15  
 Ala Arg Asp Gly Gly Phe Pro Pro Phe Ser Ser Tyr Glu Lys Leu Asp  
 20 25 30  
 Ile Thr Val Leu Asp Val Asn Asp Asn Ala Pro  
 35 40

(2) INFORMATION FOR SEQ ID NO:62:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:62:

AAGGGGATCG ATTATGAGAA GGTCAAAGAC TACACCATTG AGATTGTGGC TGTGGACTCT 60  
 GGCAACCCCC CACTCTCCAG CACTAACTCC CTCAAGGTGC AGGTGGTGGG CGTCAATGAC 120  
 AACGCACCGT G 131

(2) INFORMATION FOR SEQ ID NO:63:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 43 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:63:

Lys Gly Ile Asp Tyr Glu Lys Val Lys Asp Tyr Thr Ile Glu Ile Val  
 1 5 10 15  
 Ala Val Asp Ser Gly Asn Pro Pro Leu Ser Ser Thr Asn Ser Leu Lys  
 20 25 30  
 Val Gln Val Val Asp Val Asn Asp Asn Ala Pro  
 35 40

(ii) MOLECULE TYPE: cDNA

AAGGGTTTGG ACTACGAGAC CACACAGGCC TACCAGCTCA CGGTCAACGC CACAGATCAA 60  
GACAACACCA GGCCTCTGTC CACCCTGGCC AACTTGGCCA TCATCATCAC AGATGTCCAG 120

GACATGGACC CCATCTTCAT CAACCTGCCT TACAGCACCA ACATCTACGA GCATTCTCCT 180  
 CCGGGCACGA CCGTGCGCAT CATCACCGCC ATAGACCAGG ATCAAGGACG TCCCCGGGGC 240  
 ATTGGCTACA CCATCGTTTC AGGGAATACC AACAGCATCT TTGCCCTGGA CTACATCAGC 300  
 GGAGTGCTGA CTTGAATGG CCTGCTGGAC CGGGAGAACC CCCTGTACAG CCATGGCTTC 360  
 ATCCTGACTG TGAAGGGCAC GGAGCTGAAC GATGACCGCA CCCCATCTGA CGCTACAGTC 420  
 ACCACGACCT TCAATATCCT GGTATTGAC ATCAACGACA ACGCCCCACT 470

(2) INFORMATION FOR SEQ ID NO:67:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 156 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:67:

Lys	Gly	Leu	Asp	Tyr	Glu	Thr	Thr	Gln	Ala	Tyr	Gln	Leu	Thr	Val	Asn	1	5	10	15
Ala	Thr	Asp	Gln	Asp	Asn	Thr	Arg	Pro	Leu	Ser	Thr	Leu	Ala	Asn	Leu	20	25	30	
Ala	Ile	Ile	Ile	Thr	Asp	Val	Gln	Asp	Met	Asp	Pro	Ile	Phe	Ile	Asn	35	40	45	
Leu	Pro	Tyr	Ser	Thr	Asn	Ile	Tyr	Glu	His	Ser	Pro	Pro	Gly	Thr	Thr	50	55	60	
Val	Arg	Ile	Ile	Thr	Ala	Ile	Asp	Gln	Asp	Gln	Gly	Arg	Pro	Arg	Gly	65	70	75	80
Ile	Gly	Tyr	Thr	Ile	Val	Ser	Gly	Asn	Thr	Asn	Ser	Ile	Phe	Ala	Leu	85	90	95	
Asp	Tyr	Ile	Ser	Gly	Val	Leu	Thr	Leu	Asn	Gly	Leu	Leu	Asp	Arg	Glu	100	105	110	
Asn	Pro	Leu	Tyr	Ser	Gly	Gly	Phe	Ile	Leu	Thr	Val	Lys	Gly	Thr	Glu	115	120	125	
Leu	Asn	Asp	Asp	Arg	Thr	Pro	Ser	Asp	Ala	Thr	Val	Thr	Thr	Thr	Phe	130	135	140	
Asn	Ile	Leu	Val	Ile	Asp	Ile	Asn	Asp	Asn	Ala	Pro	145	150	155					

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(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:68:

AAGGGGGTCG	ATTACGAGGT	ACTACAGGCC	TTTGAGTTCC	ACGTGAGCGC	CACAGACCGA	60
GGCTCACCGG	GGCTCAGCAG	CCAGGCTCTG	GTGCGCGTGG	TGGTGCTGGA	CGACAATGAC	120
AACGCTCCCC	T					131

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 43 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:69:

Lys Gly Val Asp Tyr Glu Val Leu Gln Ala Phe Glu Phe His Val Ser  
1 5 10 15  
Ala Thr Asp Arg Gly Ser Pro Gly Leu Ser Ser Gln Ala Leu Val Arg  
20 25 30  
Val Val Val Leu Asp Asp Asn Asp Asn Ala Pro  
35 40

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 131 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:70:

AAGGGGCTGG ATTATGAGCA GTTCCAGACC CTACAACTGG GAGTGACCGC TAGTGACAGT 60  
GGAAACCCAC CATTAAAGAAG CAATATTTCA CTGACCCTTT TCGTGCTGGA CCAGAATGAT 120

AACGCCCCAA A

131

(2) INFORMATION FOR SEQ ID NO:71:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 43 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:71:

Lys Gly Leu Asp Tyr Glu Gln Phe Gln Thr Leu Gln Leu Gly Val Thr  
 1 5 10 15  
 Ala Ser Asp Ser Gly Asn Pro Pro Leu Arg Ser Asn Ile Ser Leu Thr  
 20 25 30  
 Leu Phe Val Leu Asp Gln Asn Asp Asn Ala Pro  
 35 40

(2) INFORMATION FOR SEQ ID NO:72:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:72:

AAGCGGGTTG ATTACGAGGA TGTCCAGAAA TACTCGCTGA GCATTAAGGC CCAGGATGGG 60  
 CGGCCCCCGC TCATCAATTC TTCAGGGGTG GTGTCTGTGC AGGTGCTGGA TGTCAACGAC 120  
 AATGCCCCGG A 131

(2) INFORMATION FOR SEQ ID NO:73:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 43 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:73:

Lys Arg Val Asp Tyr Glu Asp Val Gln Lys Tyr Ser Leu Ser Ile Lys  
 1 5 10 15

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Ala Gln Asp Gly Arg Pro Pro Leu Ile Asn Ser Ser Gly Val Val Ser  
20 25 30

Val Gln Val Leu Asp Val Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:74:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 125 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:74:

AAACCGGTAG ACTTTGAGCT ACAGCAGTTC TATGAAGTAG CTGTGGTGGC TTGGAACCTCT 60  
GAGGGATTTC ATGTCAAAAG GGCATTAAA GTGCAACTTT TAGATGACAA CGACAATGCC 120  
CCGAT 125

(2) INFORMATION FOR SEQ ID NO:75:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 41 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:75:

Lys Pro Val Asp Phe Glu Leu Gln Gln Phe Tyr Glu Val Ala Val Val  
1 5 10 15  
Ala Trp Asn Ser Glu Gly Phe His Val Lys Arg Val Ile Lys Val Gln  
20 25 30  
Leu Leu Asp Asp Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:76:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 125 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:76:

AAGGGATTAG ATTTTGAAC TTGCCCATT TACACATTGA TAATACAAGG AACTAACATG 60  
GCTGGTTTGT CCACTAATAC AACGGTTCTA GTTCACTTGC AGGATGAGAA TGATAACGCC 120  
CCAAA 125

(2) INFORMATION FOR SEQ ID NO:77:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 41 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:77:

Lys Gly Leu Asp Phe Glu Thr Leu Pro Ile Tyr Thr Leu Ile Ile Gln  
1 5 10 15  
Gly Thr Asn Met Ala Gly Leu Ser Thr Asn Thr Thr Val Leu Val His  
20 25 30  
Leu Gln Asp Glu Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:78:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 134 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:78:

AAGCGGGCGG ATTTGAGGC GATCCGGGAG TACAGTCTGA GGATCAAAGC GCAGGACGGG 60  
GGGCGGCCTC CCCTCAGCAA CACCACGGGC ATGGTCACAG TGCAGGTCGT GGACGTCAAT 120  
GACAACGCAC CCCT 134

(2) INFORMATION FOR SEQ ID NO:79:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 44 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

090573 061301



(xi) SEQUENCE DESCRIPTION: SEQ ID NO:79:

Lys Arg Ala Asp Phe Glu Ala Ile Arg Glu Tyr Ser Leu Arg Ile Lys  
 1 5 10 15  
 Ala Gln Asp Gly Gly Arg Pro Pro Leu Ser Asn Thr Thr Gly Met Val  
 20 25 30  
 Thr Val Gln Val Val Asp Val Asn Asp Asn Ala Pro  
 35 40

(2) INFORMATION FOR SEQ ID NO:80:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:80:

AAGCGGTTGG ATTACGAAAA GGCATCGGAA TATGAAATCT ATGTTCAAGC CGCTGACAAA 60  
 GGCGCTGTCC CTATGGCTGG CCATTGCAAA GTGTTGCTGG AGATCGTGGA TGTCAACGAC 120  
 AACGCCCCCT T 131

(2) INFORMATION FOR SEQ ID NO:81:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 43 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:81:

Lys Arg Leu Asp Tyr Glu Lys Ala Ser Glu Tyr Glu Ile Tyr Val Gln  
 1 5 10 15  
 Ala Ala Asp Lys Gly Ala Val Pro Met Ala Gly His Cys Lys Val Leu  
 20 25 30  
 Leu Glu Ile Val Asp Val Asn Asp Asn Ala Pro  
 35 40

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(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 131 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEO ID NO:82:

(2) INFORMATION FOR SEQ ID NO:83:

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:83:

(2) INFORMATION FOR SEQ ID NO:84:

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:84:

AAAGGGTTAG ATTTTCGAGGG CACTAAAGAT TCAGCGTTTA AAATAGTGGC AGCTGACACA 60  
GGGAAGCCCA GCCTCAACCA GACAGCCCTG GTGAGAGTAG AGCTGGAGGA TGAGAACGAC 120

AACGCCCCAA T

131

(2) INFORMATION FOR SEQ ID NO:85:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:85:

Lys	Gly	Leu	Asp	Phe	Glu	Gly	Thr	Lys	Asp	Ser	Ala	Phe	Lys	Ile	Val
1					5			10					15		
Ala	Ala	Asp	Thr	Gly	Lys	Pro	Ser	Leu	Asn	Gln	Thr	Ala	Leu	Val	Arg
			20					25					30		
Val	Glu	Leu	Glu	Asp	Glu	Asn	Asp	Asn	Ala	Pro					
			35				40								

(2) INFORMATION FOR SEQ ID NO:86:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 130 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:86:

AAGGGTGTGG	ATTTTGAAG	TGTGCGTAGC	TACAGGCTGG	TTATTCGTGC	TCAAGATGGA	60
GGCAGCCCCT	CCAGAAGTAA	CACCACCCAG	CTCTTGGTCA	ACGTCATCGA	TCGAATGACA	120
ATGCGCCGCT						130

(2) INFORMATION FOR SEQ ID NO:87:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 43 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:87:

Lys	Gly	Val	Asp	Phe	Glu	Ser	Val	Arg	Ser	Tyr	Arg	Leu	Val	Ile	Arg
1					5			10				15			

- 68 -

Ala Gln Asp Gly Gly Ser Pro Ser Arg Ser Asn Thr Thr Gln Leu Leu  
20 25 30

Val Asn Val Ile Asp Val Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:88:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 131 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:88:

AAGGGTGTGG ACTTCGAGCT GACACATCTG TATGAGATTT GGATTGAGGC TGCCGATGGA 60  
GACACGCCAA GTCTGCGTAG TGTA ACTCTT ATAACGCTCA ACGTAACGGA TGCCAATGAC 120  
AATGCTCCCA A 131

(2) INFORMATION FOR SEQ ID NO:89:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 43 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:89:

Lys Gly Val Asp Phe Glu Leu Thr His Leu Tyr Glu Ile Trp Ile Glu  
1 5 10 15  
Ala Ala Asp Gly Asp Thr Pro Ser Leu Arg Ser Val Thr Leu Ile Thr  
20 25 30  
Leu Asn Val Thr Asp Ala Asn Asp Asn Ala Pro  
35 40

(2) INFORMATION FOR SEQ ID NO:90:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 441 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

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CAAGGCGTTT	GATTTTGAAG	AGACAAGTAG	ATATGTGTTG	AGTGTGGAAG	CTAAGGATGG	60
AGGAGTACAC	ACAGCTCACT	GTAATGTTCA	AATAGAAATT	GTTGACGAGA	ATGACAATGC	120
CCCAGAGGTG	ACATTCATGT	CCTTCTCTAA	CCAGATTCCA	GAGGATTGAG	ACCTTGGAAC	180
TGTAATAGCC	CTCATAAAAG	TGCGAGACAA	GGATTCTGGG	CAAAATGGCA	TGGTGACATG	240
CTATACTCAG	GAAGAAGTTC	CTTTCAAATT	AGAATCCACC	TCGAAGAATT	ATTACAAGCT	300
GGTGATTGCT	GGAGCCCTAA	ACCGGGAGCA	GACAGCAGAC	TACAACGTCA	CAATCATAGC	360
CACCGACAAG	GGCAAACCAG	CCCTTTCCTC	CAGGACAAGC	ATCACCTGTC	ACATCTCCGA	420
CATCAACGAT	AATGCCCCCG	T				441

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 146 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:91:

[illegible]

(2) INFORMATION FOR SEQ ID NO:92:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 131 base pairs  
    (B) TYPE: nucleic acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:92:

AAGCGAGTGG ATTACGAGGC CACTCGGAAT TATAAGCTGA GAGTTAAGGC TACTGATCTT      60  
GGGATTCCAC CGAGATCTTC TAACATGACA CTGTTTCATTC ATGTCCTTGA TGTTAACGAC      120  
AACGCTCCCT T      131

(2) INFORMATION FOR SEQ ID NO:93:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 43 amino acids  
    (B) TYPE: amino acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:93:

Lys Arg Val Asp Tyr Glu Ala Thr Arg Asn Tyr Lys Leu Arg Val Lys  
1                      5                      10                      15  
Ala Thr Asp Leu Gly Ile Pro Pro Arg Ser Ser Asn Met Thr Leu Phe  
                    20                      25                      30  
Ile His Val Leu Asp Val Asn Asp Asn Ala Pro  
                    35                      40

(2) INFORMATION FOR SEQ ID NO:94:

- (i) SEQUENCE CHARACTERISTICS:  
    (A) LENGTH: 4104 base pairs  
    (B) TYPE: nucleic acid  
    (C) STRANDEDNESS: single  
    (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

- (A) NAME/KEY: CDS  
    (B) LOCATION: 495..3572

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:94:

0960573-091301

CCTCTATTTCG ACATTCTCTT TGGATTGTTT TGCTATAACT TGAAATTG GATGTCACAA	60
ACGAAACTGT CATCTGTTTC CGCCAACTG TGGTTCTGCT AATCTCCCAG GCTGGCAGCA	120
TTGGAGACTT GCTGACTTCT TTCATCCCC ACTCTTTTCA CCTGAAATTC CTTTCCTTGG	180
TTTTGCTCTA AGTCCTATGC TTCAGTCAGG GGCCAACCAA ATCTCACTGC CTCCTTTTTTA	240
TCATGAAGCC TTTGATCACT GATAGTTCTT TTTATATCTT GAAAAATCAC CCTTCCCAGT	300
ACAGTTAATA TTTAGTATCT CTACTCATCT TGGCACTTAC TCACAGCTCC ATAATTCAGT	360
CGTTTTTCGTA CCTCTTCATG GTGATGGGGA GCCCTTTGGA GGTGGTGA CTGCTTTATA	420
CTCCTCATGA TGCTTCACAT GTGGCAGGCG TGGAGTGCCC GGAGGCGGCC CTCCTGATTC	480
TGGGGCCTCC CAGG ATG GAG CCC CTG AGG CAC AGC CCA GGC CCT GGG GGG	530
Met Glu Pro Leu Arg His Ser Pro Gly Pro Gly Gly	
1 5 10	
CAA CGG CTA CTG CTG CCC TCC ATG CTG CTA GCA CTG CTG CTC CTG CTG	578
Gln Arg Leu Leu Leu Pro Ser Met Leu Leu Ala Leu Leu Leu Leu Leu	
15 20 25	
GCT CCA TCC CCA GGC CAC GCC ACT CGG GTA GTG TAC AAG GTG CCG GAG	626
Ala Pro Ser Pro Gly His Ala Thr Arg Val Val Tyr Lys Val Pro Glu	
30 35 40	
GAA CAG CCA CCC AAC ACC CTC ATT GGG AGC CTC GCA GCC GAC TAT GGT	674
Glu Gln Pro Pro Asn Thr Leu Ile Gly Ser Leu Ala Ala Asp Tyr Gly	
45 50 55 60	
TTT CCA GAT GTG GGG CAC CTG TAC AAG CTA GAG GTG GGT GCC CCG TAC	722
Phe Pro Asp Val Gly His Leu Tyr Lys Leu Glu Val Gly Ala Pro Tyr	
65 70 75	
CTT CGC GTG GAT GGC AAG ACA GGT GAC ATT TTC ACC ACC GAG ACC TCC	770
Leu Arg Val Asp Gly Lys Thr Gly Asp Ile Phe Thr Thr Glu Thr Ser	
80 85 90	
ATC GAC CGT GAG GGG CTC CGT GAA TGC CAG AAC CAG CTC CCT GGT GAT	818
Ile Asp Arg Glu Gly Leu Arg Glu Cys Gln Asn Gln Leu Pro Gly Asp	
95 100 105	
CCC TGC ATC CTG GAG TTT GAG GTA TCT ATC ACA GAC CTC GTG CAG AAT	866
Pro Cys Ile Leu Glu Phe Glu Val Ser Ile Thr Asp Leu Val Gln Asn	
110 115 120	
GCG AGC CCC CGG CTG CTA GAG GGC CAG ATA GAA GTA CAA GAC ATC AAT	914
Ala Ser Pro Arg Leu Leu Glu Gly Gln Ile Glu Val Gln Asp Ile Asn	
125 130 135 140	
GAC AAC ACA CCC AAC TTC GCC TCA CCA GTC ATC ACT CTG GCC ATC CCT	962
Asp Asn Thr Pro Asn Phe Ala Ser Pro Val Ile Thr Leu Ala Ile Pro	
145 150 155	
GAG AAC ACC AAC ATC GGC TCA CTC TTC CCC ATC CCG CTG GCT TCA GAC	1010
Glu Asn Thr Asn Ile Gly Ser Leu Phe Pro Ile Pro Leu Ala Ser Asp	
160 165 170	

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CGT GAT GCT GGT CCC AAC GGT GTG GCA TCC TAT GAG CTG CAG GTG GCA Arg Asp Ala Gly Pro Asn Gly Val Ala Ser Tyr Glu Leu Gln Val Ala 175 180 185	1058
GAG GAC CAG GAG GAG AAG CAA CCA CAG CTC ATT GTG ATG GGC AAC CTG Glu Asp Gln Glu Glu Lys Gln Pro Gln Leu Ile Val Met Gly Asn Leu 190 195 200	1106
GAC CGT GAG CGC TGG GAC TCC TAT GAC CTC ACC ATC AAG GTG CAG GAT Asp Arg Glu Arg Trp Asp Ser Tyr Asp Leu Thr Ile Lys Val Gln Asp 205 210 215 220	1154
GGC GGC AGC CCC CCA CGC GCC ACG AGT GCC CTG CTG CGT GTC ACC GTG Gly Gly Ser Pro Pro Arg Ala Thr Ser Ala Leu Leu Arg Val Thr Val 225 230 235	1202
CTT GAC ACC AAT GAC AAC GCC CCC AAG TTT GAG CGG CCC TCC TAT GAG Leu Asp Thr Asn Asp Asn Ala Pro Lys Phe Glu Arg Pro Ser Tyr Glu 240 245 250	1250
GCC GAA CTA TCT GAG AAT AGC CCC ATA GGC CAC TCG GTC ATC CAG GTG Ala Glu Leu Ser Glu Asn Ser Pro Ile Gly His Ser Val Ile Gln Val 255 260 265	1298
AAG GCC AAT GAC TCA GAC CAA GGT GCC AAT GCA GAA ATC GAA TAC ACA Lys Ala Asn Asp Ser Asp Gln Gly Ala Asn Ala Glu Ile Glu Tyr Thr 270 275 280	1346
TTC CAC CAG GCG CCC GAA GTT GTG AGG CGT CTT CTT CGA CTG GAC AGG Phe His Gln Ala Pro Glu Val Val Arg Arg Leu Leu Arg Leu Asp Arg 285 290 295 300	1394
AAC ACT GGA CTT ATC ACT GTT CAG GGC CCG GTG GAC CGT GAG GAC CTA Asn Thr Gly Leu Ile Thr Val Gln Gly Pro Val Asp Arg Glu Asp Leu 305 310 315	1442
AGC ACC CTG CGC TTC TCA GTG CTT GCT AAG GAC CGA GGC ACC AAC CCC Ser Thr Leu Arg Phe Ser Val Leu Ala Lys Asp Arg Gly Thr Asn Pro 320 325 330	1490
AAG AGT GCC CGT GCC CAG GTG GTT GTG ACC GTG AAG GAC ATG AAT GAC Lys Ser Ala Arg Ala Gln Val Val Thr Val Lys Asp Met Asn Asp 335 340 345	1538
AAT GCC CCC ACC ATT GAG ATC CGG GGC ATA GGG CTA GTG ACT CAT CAA Asn Ala Pro Thr Ile Glu Ile Arg Gly Ile Gly Leu Val Thr His Gln 350 355 360	1586
GAT GGG ATG GCT AAC ATC TCA GAG GAT GTG GCA GAG GAG ACA GCT GTG Asp Gly Met Ala Asn Ile Ser Glu Asp Val Ala Glu Glu Thr Ala Val 365 370 375 380	1634
GCC CTG GTG CAG GTG TCT GAC CGA GAT GAG GGA GAG AAT GCA GCT GTC Ala Leu Val Gln Val Ser Asp Arg Asp Glu Gly Glu Asn Ala Ala Val 385 390 395	1682
ACC TGT GTG GTG GCA GGT GAT GTG CCC TTC CAG CTG CGC CAG GCC AGT Thr Cys Val Val Ala Gly Asp Val Pro Phe Gln Leu Arg Gln Ala Ser 400 405 410	1730

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GAG Glu	ACA Thr	GGC Gly	AGT Ser	GAC Asp	AGC Ser	AAG Lys	AAG Lys	AAG Lys	TAT Tyr	TTC Phe	CTG Leu	CAG Gln	ACT Thr	ACC Thr	ACC Thr	1778
415 420 425																
CCG Pro	CTA Leu	GAC Asp	TAC Tyr	GAG Glu	AAG Lys	GTC Val	AAA Lys	GAC Asp	TAC Tyr	ACC Thr	ATT Ile	GAG Glu	ATT Ile	GTG Val	GCT Ala	1826
430 435 440																
GTG Val	GAC Asp	TCT Ser	GGC Gly	AAC Asn	CCC Pro	CCA Pro	CTC Leu	TCC Ser	AGC Ser	ACT Thr	AAC Asn	TCC Ser	CTC Leu	AAG Lys	GTG Val	1874
445 450 455 460																
CAG Gln	GTG Val	GTG Val	GAC Asp	GTC Val	AAT Asn	GAC Asp	AAC Asn	GCA Ala	CCT Pro	GTC Val	TTC Phe	ACT Thr	CAG Gln	AGT Ser	GTC Val	1922
465 470 475 480 485 490																
ACT Thr	GAG Glu	GTC Val	GCC Ala	TTC Phe	CCG Pro	GAA Glu	AAC Asn	AAC Asn	AAG Lys	CCT Pro	GGT Gly	GAA Glu	GTG Val	ATT Ile	GCT Ala	1970
480 485 490																
GAG Glu	ATC Ile	ACT Thr	GCC Ala	AGT Ser	GAT Asp	GCT Ala	GAC Asp	TCT Ser	GGC Gly	TCT Ser	AAT Asn	GCT Ala	GAG Glu	CTG Leu	GTT Val	2018
495 500 505																
TAC Tyr	TCT Ser	CTG Leu	GAG Glu	CCT Pro	GAG Glu	CCG Pro	GCT Ala	GCT Ala	AAG Lys	GGC Gly	CTC Leu	TTC Phe	ACC Thr	ATC Ile	TCA Ser	2066
510 515 520																
CCC Pro	GAG Glu	ACT Thr	GGA Gly	GAG Glu	ATC Ile	CAG Gln	GTG Val	AAG Lys	ACA Thr	TCT Ser	CTG Leu	GAT Asp	CGG Arg	GAA Glu	CAG Gln	2114
525 530 535 540																
CGG Arg	GAG Glu	AGC Ser	TAT Tyr	GAG Glu	TTG Leu	AAG Lys	GTG Val	GTG Val	GCA Ala	GCT Ala	GAC Asp	CGG Arg	GGC Gly	AGT Ser	CCT Pro	2162
545 550 555																
AGC Ser	CTC Leu	CAG Gln	GGC Gly	ACA Thr	GCC Ala	ACT Thr	GTC Val	CTT Leu	GTC Val	AAT Asn	GTG Val	CTG Leu	GAC Asp	TGC Cys	AAT Asn	2210
560 565 570 575																
GAC Asp	AAT Asn	GAC Asp	CCC Pro	AAA Lys	TTT Phe	ATG Met	CTG Leu	AGT Ser	GGC Gly	TAC Tyr	AAC Asn	TTC Phe	TCA Ser	GTG Val	ATG Met	2258
580 585																
GAG Glu	AAC Asn	ATG Met	CCA Pro	GCA Ala	CTG Leu	AGT Ser	CCA Pro	GTG Val	GGC Gly	ATG Met	GTG Val	ACT Thr	GTC Val	ATT Ile	GAT Asp	2306
590 595 600																
GGA Gly	GAC Asp	AAG Lys	GGG Gly	GAG Glu	AAT Asn	GCC Ala	CAG Gln	GTG Val	CAG Gln	CTC Leu	TCA Ser	GTG Val	GAG Glu	CAG Gln	GAC Asp	2354
605 610 615 620																
AAC Asn	GGT Gly	GAC Asp	TTT Phe	GTT Val	ATC Ile	CAG Gln	AAT Asn	GGC Gly	ACA Thr	GGC Gly	ACC Thr	ATC Ile	CTA Leu	TCC Ser	AGC Ser	2402
625 630 635																
CTG Leu	AGC Ser	TTT Phe	GAT Asp	CGA Arg	GAG Glu	CAA Gln	CAA Gln	AGC Ser	ACC Thr	TAC Tyr	ACC Thr	TTC Phe	CAG Gln	CTG Leu	AAG Lys	2450
640 645 650																

GCA GTG GAT GGT GGC GTC CCA CCT CGC TCA GCT TAC GTT GGT GTC ACC Ala Val Asp Gly Gly Val Pro Pro Arg Ser Ala Tyr Val Gly Val Thr 655 660 665	2498
ATC AAT GTG CTG GAC GAG AAT GAC AAC GCA CCC TAT ATC ACT GCC CCT Ile Asn Val Leu Asp Glu Asn Asp Asn Ala Pro Tyr Ile Thr Ala Pro 670 675 680	2546
TCT AAC ACC TCT CAC AAG CTG CTG ACC CCC CAG ACA CGT CTT GGT GAG Ser Asn Thr Ser His Lys Leu Leu Thr Pro Gln Thr Arg Leu Gly Glu 685 690 700	2594
ACG GTC AGC CAG GTG GCA GCC GAG GAC TTT GAC TCT GGT GTC AAT GCC Thr Val Ser Gln Val Ala Ala Glu Asp Phe Asp Ser Gly Val Asn Ala 705 710 715	2642
GAG CTG ATC TAC AGC ATT GCA GGT GGC AAC CCT TAT GGA CTC TTC CAG Glu Leu Ile Tyr Ser Ile Ala Gly Gly Asn Pro Tyr Gly Leu Phe Gln 720 725 730	2690
ATT GGG TCA CAT TCA GGT GCC ATC ACC CTG GAG AAG GAG ATT GAG CGG Ile Gly Ser His Ser Gly Ala Ile Thr Leu Glu Lys Glu Ile Glu Arg 735 740 745	2738
CGC CAC CAT GGG CTA CAC CGC CTG GTG GTG AAG GTC AGT GAC CGC GGC Arg His His Gly Leu His Arg Leu Val Val Lys Val Ser Asp Arg Gly 750 755 760	2786
AAG CCC CCA CGC TAT GGC ACA GCC TTG GTC CAT CTT TAT GTC AAT GAG Lys Pro Pro Arg Tyr Gly Thr Ala Leu Val His Leu Tyr Val Asn Glu 765 770 775 780	2834
ACT CTG GCC AAC CGC ACG CTG CTG GAG ACC CTC CTG GGC CAC AGC CTG Thr Leu Ala Asn Arg Thr Leu Leu Glu Thr Leu Leu Gly His Ser Leu 785 790 795	2882
GAC ACG CCG CTG GAT ATT GAC ATT GCT GGG GAT CCA GAA TAT GAG CGC Asp Thr Pro Leu Asp Ile Asp Ile Ala Gly Asp Pro Glu Tyr Glu Arg 800 805 810	2930
TCC AAG CAG CGT GGC AAC ATT CTC TTT GGT GTG GTG GCT GGT GTG GTG Ser Lys Gln Arg Gly Asn Ile Leu Phe Gly Val Val Ala Gly Val Val 815 820 825	2978
GCC GTG GCC TTG CTC ATC GCC CTG GCG GTT CTT GTG CGC TAC TGC AGA Ala Val Ala Leu Leu Ile Ala Leu Ala Val Leu Val Arg Tyr Cys Arg 830 835 840	3026
CAG CGG GAG GCC AAA AGT GGT TAC CAG GCT GGT AAG AAG GAG ACC AAG Gln Arg Glu Ala Lys Ser Gly Tyr Gln Ala Gly Lys Lys Glu Thr Lys 845 850 855 860	3074
GAC CTG TAT GCC CCC AAG CCC AGT GGC AAG GCC TCC AAG GGA AAC AAA Asp Leu Tyr Ala Pro Lys Pro Ser Gly Lys Ala Ser Lys Gly Asn Lys 865 870 875	3122
AGC AAA GGC AAG AAG AGC AAG TCC CCA AAG CCC GTG AAG CCA GTG GAG Ser Lys Gly Lys Ser Lys Ser Pro Lys Pro Val Lys Pro Val Glu 880 885 890	3170

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GAC GAG GAT GAG GCC GGG CTG CAG AAG TCC CTC AAG TTC AAC CTG ATG Asp Glu Asp Glu Ala Gly Leu Gln Lys Ser Leu Lys Phe Asn Leu Met 895 900 905	3218
AGC GAT GCC CCT GGG GAC AGT CCC CGC ATC CAC CTG CCC CTC AAC TAC Ser Asp Ala Pro Gly Asp Ser Pro Arg Ile His Leu Pro Leu Asn Tyr 910 915 920	3266
CCA CCA GGC AGC CCT GAC CTG GGC CGC CAC TAT CGC TCT AAC TCC CCA Pro Pro Gly Ser Pro Asp Leu Gly Arg His Tyr Arg Ser Asn Ser Pro 925 930 935 940	3314
CTG CCT TCC ATC CAG CTG CAG CCC CAG TCA CCC TCA GCC TCC AAG AAG Leu Pro Ser Ile Gln Leu Gln Pro Gln Ser Pro Ser Ala Ser Lys Lys 945 950 955	3362
CAC CAG GTG GTA CAG GAC CTG CCA CCT GCA AAC ACA TTC GTG GGC ACC His Gln Val Val Gln Asp Leu Pro Pro Ala Asn Thr Phe Val Gly Thr 960 965 970	3410
GGG GAC ACC ACG TCC ACG GGC TCT GAG CAG TAC TCC GAC TAC AGC TAC Gly Asp Thr Thr Ser Thr Gly Ser Glu Gln Tyr Ser Asp Tyr Ser Tyr 975 980 985	3458
CGC ACC AAC CCC CCC AAA TAC CCC AGC AAG CAG GTA GGC CAG CCC TTT Arg Thr Asn Pro Pro Lys Tyr Pro Ser Lys Gln Val Gly Gln Pro Phe 990 995 1000	3506
CAG CTC AGC ACA CCC CAG CCC CTA CCC CAC CCC TAC CAC GGA GCC ATC Gln Leu Ser Thr Pro Gln Pro Leu Pro His Pro Tyr His Gly Ala Ile 1005 1010 1015 1020	3554
TGG ACC GAG GTG TGG GAG TGATGGAGCA GGTTTACTGT GCCTGCCCCGT Trp Thr Glu Val Trp Glu 1025	3602
GTTGGGGGCC AGCCTGAGCC AGCAGTGGGA GGTGGGGCCT TAGTGCCTCA CCGGGCACAC	3662
GGATTAGGCT GAGTGAAGAT TAAGGGAGGG TGTGCTCTGT GGTCTCCTCC CTGCCCTCTC	3722
CCCACTGGGG AGAGACCTGT GATTTGCCAA GTCCCTGGAC CCTGGACCAG CTACTGGGCC	3782
TTATGGGTTG GGGGTGGTAG GCAGGTGAGC GTAAGTGGGG AGGGAAATGG GTAAGAAGTC	3842
TACTCCAAAC CTAGGTCTCT ATGTCAGACC AGACCTAGGT GCTTCTCTAG GAGGGAAACA	3902
GGGAGACCTG GGGTCCTGTG GATAACTGAG TGGGGAGTCT GCCAGGGGAG GGCACCTTCC	3962
CATTGTGCCT TCTGTGTGTA TTGTGCATTA ACCTCTTCCT CACCACTAGG CTTCTGGGGC	4022
TGGGTCCCAC ATGCCCTTGA CCCTGACAAT AAAGTTCTCT ATTTTGGAA AAAAAAAAAA	4082
AAAAAAAAAA AAAAAAAAAA AA	4104

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(2) INFORMATION FOR SEQ ID NO:95:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1026 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:95:

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Met Glu Pro Leu Arg His Ser Pro Gly Pro Gly Gly Gln Arg Leu Leu
 1          5          10          15
Leu Pro Ser Met Leu Leu Ala Leu Leu Leu Leu Ala Pro Ser Pro
          20          25          30
Gly His Ala Thr Arg Val Val Tyr Lys Val Pro Glu Glu Gln Pro Pro
          35          40          45
Asn Thr Leu Ile Gly Ser Leu Ala Ala Asp Tyr Gly Phe Pro Asp Val
          50          55          60
Gly His Leu Tyr Lys Leu Glu Val Gly Ala Pro Tyr Leu Arg Val Asp
          65          70          75          80
Gly Lys Thr Gly Asp Ile Phe Thr Thr Glu Thr Ser Ile Asp Arg Glu
          85          90          95
Gly Leu Arg Glu Cys Gln Asn Gln Leu Pro Gly Asp Pro Cys Ile Leu
          100          105          110
Glu Phe Glu Val Ser Ile Thr Asp Leu Val Gln Asn Ala Ser Pro Arg
          115          120          125
Leu Leu Glu Gly Gln Ile Glu Val Gln Asp Ile Asn Asp Asn Thr Pro
          130          135          140
Asn Phe Ala Ser Pro Val Ile Thr Leu Ala Ile Pro Glu Asn Thr Asn
          145          150          155          160
Ile Gly Ser Leu Phe Pro Ile Pro Leu Ala Ser Asp Arg Asp Ala Gly
          165          170          175
Pro Asn Gly Val Ala Ser Tyr Glu Leu Gln Val Ala Glu Asp Gln Glu
          180          185          190
Glu Lys Gln Pro Gln Leu Ile Val Met Gly Asn Leu Asp Arg Glu Arg
          195          200          205
Trp Asp Ser Tyr Asp Leu Thr Ile Lys Val Gln Asp Gly Gly Ser Pro
          210          215          220
Pro Arg Ala Thr Ser Ala Leu Leu Arg Val Thr Val Leu Asp Thr Asn
          225          230          235          240
Asp Asn Ala Pro Lys Phe Glu Arg Pro Ser Tyr Glu Ala Glu Leu Ser
          245          250          255
Glu Asn Ser Pro Ile Gly His Ser Val Ile Gln Val Lys Ala Asn Asp
          260          265          270

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09660573 "061307"

Ser Asp Gln Gly Ala Asn Ala Glu Ile Glu Tyr Thr Phe His Gln Ala  
275 280 285

Pro Glu Val Val Arg Arg Leu Leu Arg Leu Asp Arg Asn Thr Gly Leu  
290 295 300

Ile Thr Val Gln Gly Pro Val Asp Arg Glu Asp Leu Ser Thr Leu Arg  
305 310 315 320

Phe Ser Val Leu Ala Lys Asp Arg Gly Thr Asn Pro Lys Ser Ala Arg  
325 330 335

Ala Gln Val Val Val Thr Val Lys Asp Met Asn Asp Asn Ala Pro Thr  
340 345 350

Ile Glu Ile Arg Gly Ile Gly Leu Val Thr His Gln Asp Gly Met Ala  
355 360 365

Asn Ile Ser Glu Asp Val Ala Glu Glu Thr Ala Val Ala Leu Val Gln  
370 375 380

Val Ser Asp Arg Asp Glu Gly Glu Asn Ala Ala Val Thr Cys Val Val  
385 390 395 400

Ala Gly Asp Val Pro Phe Gln Leu Arg Gln Ala Ser Glu Thr Gly Ser  
405 410 415

Asp Ser Lys Lys Lys Tyr Phe Leu Gln Thr Thr Thr Pro Leu Asp Tyr  
420 425 430

Glu Lys Val Lys Asp Tyr Thr Ile Glu Ile Val Ala Val Asp Ser Gly  
435 440 445

Asn Pro Pro Leu Ser Ser Thr Asn Ser Leu Lys Val Gln Val Val Asp  
450 455 460

Val Asn Asp Asn Ala Pro Val Phe Thr Gln Ser Val Thr Glu Val Ala  
465 470 475 480

Phe Pro Glu Asn Asn Lys Pro Gly Glu Val Ile Ala Glu Ile Thr Ala  
485 490 495

Ser Asp Ala Asp Ser Gly Ser Asn Ala Glu Leu Val Tyr Ser Leu Glu  
500 505 510

Pro Glu Pro Ala Ala Lys Gly Leu Phe Thr Ile Ser Pro Glu Thr Gly  
515 520 525

Glu Ile Gln Val Lys Thr Ser Leu Asp Arg Glu Gln Arg Glu Ser Tyr  
530 535 540

Glu Leu Lys Val Val Ala Ala Asp Arg Gly Ser Pro Ser Leu Gln Gly  
545 550 555 560

Thr Ala Thr Val Leu Val Asn Val Leu Asp Cys Asn Asp Asn Asp Pro  
565 570 575

Lys Phe Met Leu Ser Gly Tyr Asn Phe Ser Val Met Glu Asn Met Pro  
580 585 590

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Ala Leu Ser Pro Val Gly Met Val Thr Val Ile Asp Gly Asp Lys Gly  
595 600 605

Glu Asn Ala Gln Val Gln Leu Ser Val Glu Gln Asp Asn Gly Asp Phe  
610 615 620

Val Ile Gln Asn Gly Thr Gly Thr Ile Leu Ser Ser Leu Ser Phe Asp  
625 630 635 640

Arg Glu Gln Gln Ser Thr Tyr Thr Phe Gln Leu Lys Ala Val Asp Gly  
645 650 655

Gly Val Pro Pro Arg Ser Ala Tyr Val Gly Val Thr Ile Asn Val Leu  
660 665 670

Asp Glu Asn Asp Asn Ala Pro Tyr Ile Thr Ala Pro Ser Asn Thr Ser  
675 680 685

His Lys Leu Leu Thr Pro Gln Thr Arg Leu Gly Glu Thr Val Ser Gln  
690 695 700

Val Ala Ala Glu Asp Phe Asp Ser Gly Val Asn Ala Glu Leu Ile Tyr  
705 710 715 720

Ser Ile Ala Gly Gly Asn Pro Tyr Gly Leu Phe Gln Ile Gly Ser His  
725 730 735

Ser Gly Ala Ile Thr Leu Glu Lys Glu Ile Glu Arg Arg His His Gly  
740 745 750

Leu His Arg Leu Val Val Lys Val Ser Asp Arg Gly Lys Pro Pro Arg  
755 760 765

Tyr Gly Thr Ala Leu Val His Leu Tyr Val Asn Glu Thr Leu Ala Asn  
770 775 780

Arg Thr Leu Leu Glu Thr Leu Leu Gly His Ser Leu Asp Thr Pro Leu  
785 790 795 800

Asp Ile Asp Ile Ala Gly Asp Pro Glu Tyr Glu Arg Ser Lys Gln Arg  
805 810 815

Gly Asn Ile Leu Phe Gly Val Val Ala Gly Val Val Ala Val Ala Leu  
820 825 830

Leu Ile Ala Leu Ala Val Leu Val Arg Tyr Cys Arg Gln Arg Glu Ala  
835 840 845

Lys Ser Gly Tyr Gln Ala Gly Lys Lys Glu Thr Lys Asp Leu Tyr Ala  
850 855 860

Pro Lys Pro Ser Gly Lys Ala Ser Lys Gly Asn Lys Ser Lys Gly Lys  
865 870 875 880

Lys Ser Lys Ser Pro Lys Pro Val Lys Pro Val Glu Asp Glu Asp Glu  
885 890 895

Ala Gly Leu Gln Lys Ser Leu Lys Phe Asn Leu Met Ser Asp Ala Pro  
900 905 910

00660573 064300

Gly Asp Ser Pro Arg Ile His Leu Pro Leu Asn Tyr Pro Pro Gly Ser  
 915 920 925

Pro Asp Leu Gly Arg His Tyr Arg Ser Asn Ser Pro Leu Pro Ser Ile  
 930 935 940

Gln Leu Gln Pro Gln Ser Pro Ser Ala Ser Lys Lys His Gln Val Val  
 945 950 955 960

Gln Asp Leu Pro Pro Ala Asn Thr Phe Val Gly Thr Gly Asp Thr Thr  
 965 970 975

Ser Thr Gly Ser Glu Gln Tyr Ser Asp Tyr Ser Tyr Arg Thr Asn Pro  
 980 985 990

Pro Lys Tyr Pro Ser Lys Gln Val Gly Gln Pro Phe Gln Leu Ser Thr  
 995 1000 1005

Pro Gln Pro Leu Pro His Pro Tyr His Gly Ala Ile Trp Thr Glu Val  
 1010 1015 1020

Trp Glu  
 1025

(2) INFORMATION FOR SEQ ID NO:96:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 4705 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (ix) FEATURE:  
 (A) NAME/KEY: CDS  
 (B) LOCATION: 115..2827

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:96:

CGAAAGCCAT GTCGGACTCG TCGCCCAGCG CCCAAGCGCT AACCCGCTGA AAGTTTCTCA	60
GCGAAATCTC AGGGACGATC TGGACCCCGC TGAGAGGAAC TGCTTTTGAG TGAG ATG	117
	Met
	1
GTC CCA GAG GCC TGG AGG AGC GGA CTG GTA AGC ACC GGG AGG GTA GTG	165
Val Pro Glu Ala Trp Arg Ser Gly Leu Val Ser Thr Gly Arg Val Val	
	5 10 15
GGA GTT TTG CTT CTG CTT GGT GCC TTG AAC AAG GCT TCC ACG GTC ATT	213
Gly Val Leu Leu Leu Leu Gly Ala Leu Asn Lys Ala Ser Thr Val Ile	
	20 25 30
CAC TAT GAG ATC CCG GAG GAA AGA GAG AAG GGT TTC GCT GTG GGC AAC	261
His Tyr Glu Ile Pro Glu Glu Arg Glu Lys Gly Phe Ala Val Gly Asn	
	35 40 45

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GTG	GTC	GCG	AAC	CTT	GGT	TTG	GAT	CTC	GGT	AGC	CTC	TCA	GCC	CGC	AGG	309
Val	Val	Ala	Asn	Leu	Gly	Leu	Asp	Leu	Gly	Ser	Leu	Ser	Ala	Arg	Arg	
50					55					60					65	
TTC	CCG	GTG	GTG	TCT	GGA	GCT	AGC	CGA	AGA	TTC	TTT	GAG	GTG	AAC	CGG	357
Phe	Pro	Val	Val	Ser	Gly	Ala	Ser	Arg	Arg	Phe	Phe	Glu	Val	Asn	Arg	
				70					75					80		
GAG	ACC	GGA	GAG	ATG	TTT	GTG	AAC	GAC	CGT	CTG	GAT	CGA	GAG	GAG	CTG	405
Glu	Thr	Gly	Glu	Met	Phe	Val	Asn	Asp	Arg	Leu	Asp	Arg	Glu	Glu	Leu	
			85					90					95			
TGT	GGG	ACA	CTG	CCC	TCT	TGC	ACT	GTA	ACT	CTG	GAG	TTG	GTA	GTG	GAG	453
Cys	Gly	Thr	Leu	Pro	Ser	Cys	Thr	Val	Thr	Leu	Glu	Leu	Val	Val	Glu	
	100						105					110				
AAC	CCG	CTG	GAG	CTG	TTC	AGC	GTG	GAA	GTG	GTG	ATC	CAG	GAC	ATC	AAC	501
Asn	Pro	Leu	Glu	Leu	Phe	Ser	Val	Glu	Val	Val	Ile	Gln	Asp	Ile	Asn	
	115					120					125					
GAC	AAC	AAT	CCT	GCT	TTC	CCT	ACC	CAG	GAA	ATG	AAA	TTG	GAG	ATT	AGC	549
Asp	Asn	Asn	Pro	Ala	Phe	Pro	Thr	Gln	Glu	Met	Lys	Leu	Glu	Ile	Ser	
130					135					140					145	
GAG	GCC	GTG	GCT	CCG	GGG	ACG	CGC	TTT	CCG	CTC	GAG	AGC	GCG	CAC	GAT	597
Glu	Ala	Val	Ala	Pro	Gly	Thr	Arg	Phe	Pro	Leu	Glu	Ser	Ala	His	Asp	
				150					155					160		
CCC	GAT	CTG	GGA	AGC	AAC	TCT	TTA	CAA	ACC	TAT	GAG	CTG	AGC	CGA	AAT	645
Pro	Asp	Leu	Gly	Ser	Asn	Ser	Leu	Gln	Thr	Tyr	Glu	Leu	Ser	Arg	Asn	
			165					170					175			
GAA	TAC	TTT	GCG	CTT	CGC	GTG	CAG	ACG	CGG	GAG	GAC	AGC	ACC	AAG	TAC	693
Glu	Tyr	Phe	Ala	Leu	Arg	Val	Gln	Thr	Arg	Glu	Asp	Ser	Thr	Lys	Tyr	
		180					185					190				
GCG	GAG	CTG	GTG	TTG	GAG	CGC	GCC	CTG	GAC	CGA	GAA	CGG	GAG	CCT	AGT	741
Ala	Glu	Leu	Val	Leu	Glu	Arg	Ala	Leu	Asp	Arg	Glu	Arg	Glu	Pro	Ser	
	195					200					205					
CTC	CAG	TTA	GTG	CTG	ACG	GCG	TTG	GAC	GGA	GGG	ACC	CCA	GCT	CTC	TCC	789
Leu	Gln	Leu	Val	Leu	Thr	Ala	Leu	Asp	Gly	Gly	Thr	Pro	Ala	Leu	Ser	
210					215					220					225	
GCC	AGC	CTG	CCT	ATT	CAC	ATC	AAG	GTG	CTG	GAC	GCG	AAT	GAC	AAT	GCG	837
Ala	Ser	Leu	Pro	Ile	His	Ile	Lys	Val	Leu	Asp	Ala	Asn	Asp	Asn	Ala	
				230					235					240		
CCT	GTC	TTT	AAC	CAG	TCC	TTG	TAC	CGG	GCG	CGC	GTT	CCT	GGA	GGA	TGC	885
Pro	Val	Phe	Asn	Gln	Ser	Leu	Tyr	Arg	Ala	Arg	Val	Pro	Gly	Gly	Cys	
			245					250					255			
ACC	TCC	GGC	ACG	CGC	GTG	GTA	CAA	GTC	CTT	GCA	ACG	GAT	CTG	GAT	GAA	933
Thr	Ser	Gly	Thr	Arg	Val	Val	Gln	Val	Leu	Ala	Thr	Asp	Leu	Asp	Glu	
		260					265					270				
GGC	CCC	AAC	GGT	GAA	ATT	ATT	TAC	TCC	TTC	GGC	AGC	CAC	AAC	CGC	GCC	981
Gly	Pro	Asn	Gly	Glu	Ile	Ile	Tyr	Ser	Phe	Gly	Ser	His	Asn	Arg	Ala	
	275					280					285					

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GGC GTG CGG CAA CTA TTC GCC TTA GAC CTT GTA ACC GGG ATG CTG ACA Gly Val Arg Gln Leu Phe Ala Leu Asp Leu Val Thr Gly Met Leu Thr 290 295 300 305	1029
ATC AAG GGT CGG CTG GAC TTC GAG GAC ACC AAA CTC CAT GAG ATT TAC Ile Lys Gly Arg Leu Asp Phe Glu Asp Thr Lys Leu His Glu Ile Tyr 310 315 320	1077
ATC CAG GCC AAA GAC AAG GGC GCC AAT CCC GAA GGA GCA CAT TGC AAA Ile Gln Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys 325 330 335	1125
GTG TTG GTG GAG GTT GTG GAT GTG AAT GAC AAC GCC CCG GAG ATC ACA Val Leu Val Glu Val Val Asp Val Asn Asp Asn Ala Pro Glu Ile Thr 340 345 350	1173
GTC ACC TCC GTG TAC AGC CCA GTA CCC GAG GAT GCC TCT GGG ACT GTC Val Thr Ser Val Tyr Ser Pro Val Pro Glu Asp Ala Ser Gly Thr Val 355 360 365	1221
ATC GCT TTG CTC AGT GTG ACT GAC CTG GAT GCT GGC GAG AAC GGG CTG Ile Ala Leu Leu Ser Val Thr Asp Leu Asp Ala Gly Glu Asn Gly Leu 370 375 380 385	1269
GTG ACC TGC GAA GTT CCA CCG GGT CTC CCT TTC AGC CTT ACT TCT TCC Val Thr Cys Glu Val Pro Pro Gly Leu Pro Phe Ser Leu Thr Ser Ser 390 395 400	1317
CTC AAG AAT TAC TTC ACT TTG AAA ACC AGT GCA GAC CTG GAT CCG GAG Leu Lys Asn Tyr Phe Thr Leu Lys Thr Ser Ala Asp Leu Asp Arg Glu 405 410 415	1365
ACT GTG CCA GAA TAC AAC CTC AGC ATC ACC GCC CGA GAC GCC GGA ACC Thr Val Pro Glu Tyr Asn Leu Ser Ile Thr Ala Arg Asp Ala Gly Thr 420 425 430	1413
CCT TCC CTC TCA GCC CTT ACA ATA GTG CGT GTT CAA GTG TCC GAC ATC Pro Ser Leu Ser Ala Leu Thr Ile Val Arg Val Gln Val Ser Asp Ile 435 440 445	1461
AAT GAC AAC CCT CCA CAA TCT TCT CAA TCT TCC TAC GAC GTT TAC ATT Asn Asp Asn Pro Pro Gln Ser Ser Gln Ser Ser Tyr Asp Val Tyr Ile 450 455 460 465	1509
GAA GAA AAC AAC CTC CCC GGG GCT CCA ATA CTA AAC CTA AGT GTC TGG Glu Glu Asn Asn Leu Pro Gly Ala Pro Ile Leu Asn Leu Ser Val Trp 470 475 480	1557
GAC CCC GAC GCC CCG CAG AAT GCT CGG CTT TCT TTC TTT CTC TTG GAG Asp Pro Asp Ala Pro Gln Asn Ala Arg Leu Ser Phe Phe Leu Glu 485 490 495	1605
CAA GGA GCT GAA ACC GGG CTA GTG GGT CGC TAT TTC ACA ATA AAT CGT Gln Gly Ala Glu Thr Gly Leu Val Gly Arg Tyr Phe Ile Asn Arg 500 505 510	1653
GAC AAT GGC ATA GTG TCA TCC TTA GTG CCC CTA GAC TAT GAG GAT CGG Asp Asn Gly Ile Val Ser Ser Leu Val Pro Leu Asp Tyr Glu Asp Arg 515 520 525	1701

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CGG GAA TTT GAA TTA ACA GCT CAT ATC AGC GAT GGG GGC ACC CCG GTC Arg Glu Phe Glu Leu Thr Ala His Ile Ser Asp Gly Gly Thr Pro Val 530 535 540 545	1749
CTA GCC ACC AAC ATC AGC GTG AAC ATA TTT GTC ACT GAT CGC AAT GAC Leu Ala Thr Asn Ile Ser Val Asn Ile Phe Val Thr Asp Arg Asn Asp 550 555 560	1797
AAT GCC CCC CAG GTC CTA TAT CCT CGG CCA GGT GGG AGC TCG GTG GAG Asn Ala Pro Gln Val Leu Tyr Pro Arg Pro Gly Gly Ser Ser Val Glu 565 570 575	1845
ATG CTG CCT CGA GGT ACC TCA GCT GGC CAC CTA GTG TCA CGG GTG GTA Met Leu Pro Arg Gly Thr Ser Ala Gly His Leu Val Ser Arg Val Val 580 585 590	1893
GGC TGG GAC GCG GAT GCA GGG CAC AAT GCC TGG CTC TCC TAC AGT CTC Gly Trp Asp Ala Asp Ala Gly His Asn Ala Trp Leu Ser Tyr Ser Leu 595 600 605	1941
TTT GGA TCC CCT AAC CAG AGC CTT TTT GCC ATA GGG CTG CAC ACT GGT Phe Gly Ser Pro Asn Gln Ser Leu Phe Ala Ile Gly Leu His Thr Gly 610 615 620 625	1989
CAA ATC AGT ACT GCC CGT CCA GTC CAA GAC ACA GAT TCA CCC AGG CAG Gln Ile Ser Thr Ala Arg Pro Val Gln Asp Thr Asp Ser Pro Arg Gln 630 635 640	2037
ACT CTC ACT GTC TTG ATC AAA GAC AAT GGG GAG CCT TCG CTC TCC ACC Thr Leu Thr Val Leu Ile Lys Asp Asn Gly Glu Pro Ser Leu Ser Thr 645 650 655	2085
ACT GCT ACC CTC ACT GTG TCA GTA ACC GAG GAC TCT CCT GAA GCC CGA Thr Ala Thr Leu Thr Val Ser Val Thr Glu Asp Ser Pro Glu Ala Arg 660 665 670	2133
GCC GAG TTC CCC TCT GGC TCT GCC CCC CGG GAG CAG AAA AAA AAT CTC Ala Glu Phe Pro Ser Gly Ser Ala Pro Arg Glu Gln Lys Lys Asn Leu 675 680 685	2181
ACC TTT TAT CTA CTT CTT TCT CTA ATC CTG GTT TCT GTG GGC TTC GTG Thr Phe Tyr Leu Leu Leu Ser Leu Ile Leu Val Ser Val Gly Phe Val 690 695 700 705	2229
GTC ACA GTG TTC GGA GTA ATC ATA TTC AAA GTT TAC AAG TGG AAG CAG Val Thr Val Phe Gly Val Ile Ile Phe Lys Val Tyr Lys Trp Lys Gln 710 715 720	2277
TCT AGA GAC CTA TAC CGA GCC CCG GTG AGC TCA CTG TAC CGA ACA CCA Ser Arg Asp Leu Tyr Arg Ala Pro Val Ser Ser Leu Tyr Arg Thr Pro 725 730 735	2325
GGG CCC TCC TTG CAC GCG GAC GCC GTG CGG GGA GGC CTG ATG TCG CCG Gly Pro Ser Leu His Ala Asp Ala Val Arg Gly Gly Leu Met Ser Pro 740 745 750	2373
CAC CTT TAC CAT CAG GTG TAT CTC ACC ACG GAC TCC CGC CGC AGC GAC His Leu Tyr His Gln Val Tyr Leu Thr Thr Asp Ser Arg Arg Ser Asp 755 760 765	2421

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CCG CTG CTG AAG AAA CCT GGT GCA GCC AGT CCA CTG GCC AGC CGC CAG Pro Leu Leu Lys Lys Pro Gly Ala Ala Ser Pro Leu Ala Ser Arg Gln 770 775 780 785	2469
AAC ACG CTG CGG AGC TGT GAT CCG GTG TTC TAT AGG CAG GTG TTG GGT Asn Thr Leu Arg Ser Cys Asp Pro Val Phe Tyr Arg Gln Val Leu Gly 790 795 800	2517
GCA GAG AGC GCC CCT CCC GGA CAG CAA GCC CCG CCC AAC ACG GAC TGG Ala Glu Ser Ala Pro Pro Gly Gln Gln Ala Pro Pro Asn Thr Asp Trp 805 810 815	2565
CGT TTC TCT CAG GCC CAG AGA CCC GGC ACC AGC GGC TCC CAA AAT GGC Arg Phe Ser Gln Ala Gln Arg Pro Gly Thr Ser Gly Ser Gln Asn Gly 820 825 830	2613
GAT GAC ACC GGC ACC TGG CCC AAC AAC CAG TTT GAC ACA GAG ATG CTG Asp Asp Thr Gly Thr Trp Pro Asn Asn Gln Phe Asp Thr Glu Met Leu 835 840 845	2661
CAA GCC ATG ATC TTG GCG TCC GCC AGT GAA GCT GCT GAT GGG AGC TCC Gln Ala Met Ile Leu Ala Ser Ala Ser Glu Ala Ala Asp Gly Ser Ser 850 855 860 865	2709
ACC CTG GGA GGG GGT GCC GGC ACC ATG GGA TTG AGC GCC CGC TAC GGA Thr Leu Gly Gly Gly Ala Gly Thr Met Gly Leu Ser Ala Arg Tyr Gly 870 875 880	2757
CCC CAG TTC ACC CTG CAG CAC GTG CCC GAC TAC CGC CAG AAT GTC TAC Pro Gln Phe Thr Leu Gln His Val Pro Asp Tyr Arg Gln Asn Val Tyr 885 890 895	2805
ATC CCA GGC AGC AAT GCA CAC T GACCAACGCA GCTGGCAAGC GGATGGCAAG Ile Pro Gly Ser Asn Ala His 900	2857
GCCCAGCAGG TGGCAATGGC AACAAGAAGA AGTCGGCAAG AAGGAGAAGA AGTAACATGG	2917
AGGCCAGGCC AAGAGCCACA GGGCAGCCTC TCCCCGAACC AGCCCAGCTT CTCCTTACCT	2977
GCACCCAGGC CTCAGAGTTT CAGGGCTAAC CCCCAGAATA CTGGTAGGGG CCAAGGCATC	3037
TCCCTTGGA ACAGAAACAA GTGCCATCAC ACCATCCCTT CCCCAGGTGT AATATCCAAA	3097
GCAGTTCCGC TGGGAACCCC ATCCAATCAG TGGCTGTACC CATTGTTGGTA GTGGGGTTCA	3157
TGTAGACACC AAGAACCATT TGCCACACCC CGTTTAGTTA CAGCTGAACC CTCCATCTTC	3217
CAAATCAATC AGGCCCATCC ATCCCATGCC TCCCTCCTCC CCACCCCACT CCAACAGTTC	3277
CTCTTTCCCG AGTAAGGTGG TTGGGGTGTT GAAGTACCAA GTAACCTACA AGCCTCCTAG	3337
TTCTGAAAAG TTGGAAGGGC ATCATGACCT CTTGGCCTCT CCTTTGATTC TCAATCTTCC	3397
CCCAAAGCAT GGTTTGGTGC CAGCCCCTTC ACCTCCTTCC AGAGCCCAAG ATCAATGCTC	3457
AAGTTTTGGA GGACATGATC ACCATCCCCA TGGTACTGAT GCTTGCTGGA TTTAGGGAGG	3517
GCATTTTGCT ACCAAGCCTC TTCCCAACGC CCTGGGACCA GTCTTCTGTT TTGTTTTTCA	3577
TTGTTTGAGC TTTCCACTGC ATGCCTTGAC TTCCCCCACC TCCTCCTCAA ACAAGAGACT	3637

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CCACTGCATG TTCCAAGACA GTATGGGGTG GTAAGATAAG GAAGGGAAGT GTGTGGATGT	3697
GGATGGTGGG GGCATGGACA AAGCTTGACA CATCAAGTTA TCAAGGCCTT GGAGGAGGCT	3757
CTGTATGTCC TCAGGGGACT GACAACATCC TCCAGATTCC AGCCATAAAC CAATAACTAG	3817
GCTGGACCCT TCCCACTACA TAATAGGGCT CAGCCAGGCA GCCAGCTTTG GGCTGAGCTA	3877
ACAGGACCAA TGGATTAAGT GGCATTTTCAG TCCAAGGAAG CTCGAAGCAG GTTTAGGACC	3937
AGGTCCCCTT GAGAGGTCAG AGGGGCCCTCT GTGGGTGCTG GGTACTCCAG AGGTGCCACT	3997
GGTGGAAGGG TCAGCGGAGC CCCAGCAGGA AGGGTGGGCC AGCCAGGCCA TTCTTAGTCC	4057
CTGGGTGGG GAGGCAGGGA GCTAGGGCAG GGACCAAATG AACAGAAAGT CTCAGCCCAG	4117
GATGGGGCTT CTTCAACAGG CCCCTGCCCT CCTGAAGCCT CAGTCCTTCA CCTTGCCAGG	4177
TGCCGTTTCT CTTCCGTGAA GGCCACTGCC CAGGTCCCCA GTGCGCCCCC TAGTGCCCAT	4237
AGCCTGGTTA AAGTTCCCCA GTGCCTCCTT GTGATAGACC TTCTTCTCCC ACCCCCTTCT	4297
GCCCCTGGGT CCCCggccat CCAGCGGGGC TGCCAGAGAA CCCCAGACCT GCCCTTACAG	4357
TAGTGTAGCG CCCCTCCCT CTTTCGGCTG GTGTAGAATA GCCAGTAGTG TAGTGCGGTG	4417
TGCTTTTACG TGATGGCGGG TGGGCAGCGG GCGGCGGCGT CCGCGCAGCC GTCTGTCCTT	4477
GATCTGCCCC CGGCGGCCCC TGTGTGTTT TGTGCTGTGT CCAGCGCTAA GGCGACCCCC	4537
TCCCCCGTAC TGACTTCTCC TATAAGCGCT TCTCTTCGCA TAGTCACGTA GCTCCCACCC	4597
CACCCTCTTC CTGTGTCTCA CGCAAGTTTT ATACTCTAAT ATTTATATGG CTTTTTTTCT	4657
TCGACAAAAA AATAATAAAA CGTTTCTTCT GAAAAAAAAA AAAAAAAA	4705

(2) INFORMATION FOR SEQ ID NO:97:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 904 amino acids  
 (B) TYPE: amino acid  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:97:

Met	Val	Pro	Glu	Ala	Trp	Arg	Ser	Gly	Leu	Val	Ser	Thr	Gly	Arg	Val
1				5					10					15	
Val	Gly	Val	Leu	Leu	Leu	Leu	Gly	Ala	Leu	Asn	Lys	Ala	Ser	Thr	Val
			20					25					30		
Ile	His	Tyr	Glu	Ile	Pro	Glu	Glu	Arg	Glu	Lys	Gly	Phe	Ala	Val	Gly
			35				40					45			
Asn	Val	Val	Ala	Asn	Leu	Gly	Leu	Asp	Leu	Gly	Ser	Leu	Ser	Ala	Arg
	50					55					60				

Arg Phe Pro Val Val Ser Gly Ala Ser Arg Arg Phe Phe Glu Val Asn  
65 70 75 80

Arg Glu Thr Gly Glu Met Phe Val Asn Asp Arg Leu Asp Arg Glu Glu  
85 90 95

Leu Cys Gly Thr Leu Pro Ser Cys Thr Val Thr Leu Glu Leu Val Val  
100 105 110

Glu Asn Pro Leu Glu Leu Phe Ser Val Glu Val Val Ile Gln Asp Ile  
115 120 125

Asn Asp Asn Asn Pro Ala Phe Pro Thr Gln Glu Met Lys Leu Glu Ile  
130 135 140

Ser Glu Ala Val Ala Pro Gly Thr Arg Phe Pro Leu Glu Ser Ala His  
145 150 155 160

Asp Pro Asp Leu Gly Ser Asn Ser Leu Gln Thr Tyr Glu Leu Ser Arg  
165 170 175

Asn Glu Tyr Phe Ala Leu Arg Val Gln Thr Arg Glu Asp Ser Thr Lys  
180 185 190

Tyr Ala Glu Leu Val Leu Glu Arg Ala Leu Asp Arg Glu Arg Glu Pro  
195 200 205

Ser Leu Gln Leu Val Leu Thr Ala Leu Asp Gly Gly Thr Pro Ala Leu  
210 215 220

Ser Ala Ser Leu Pro Ile His Ile Lys Val Leu Asp Ala Asn Asp Asn  
225 230 235 240

Ala Pro Val Phe Asn Gln Ser Leu Tyr Arg Ala Arg Val Pro Gly Gly  
245 250 255

Cys Thr Ser Gly Thr Arg Val Val Gln Val Leu Ala Thr Asp Leu Asp  
260 265 270

Glu Gly Pro Asn Gly Glu Ile Ile Tyr Ser Phe Gly Ser His Asn Arg  
275 280 285

Ala Gly Val Arg Gln Leu Phe Ala Leu Asp Leu Val Thr Gly Met Leu  
290 295 300

Thr Ile Lys Gly Arg Leu Asp Phe Glu Asp Thr Lys Leu His Glu Ile  
305 310 315 320

Tyr Ile Gln Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys  
325 330 335

Lys Val Leu Val Glu Val Val Asp Val Asn Asp Asn Ala Pro Glu Ile  
340 345 350

Thr Val Thr Ser Val Tyr Ser Pro Val Pro Glu Asp Ala Ser Gly Thr  
355 360 365

Val Ile Ala Leu Leu Ser Val Thr Asp Leu Asp Ala Gly Glu Asn Gly  
370 375 380

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Leu Val Thr Cys Glu Val Pro Pro Gly Leu Pro Phe Ser Leu Thr Ser  
 385 390 395 400  
 Ser Leu Lys Asn Tyr Phe Thr Leu Lys Thr Ser Ala Asp Leu Asp Arg  
 405 410 415  
 Glu Thr Val Pro Glu Tyr Asn Leu Ser Ile Thr Ala Arg Asp Ala Gly  
 420 425 430  
 Thr Pro Ser Leu Ser Ala Leu Thr Ile Val Arg Val Gln Val Ser Asp  
 435 440 445  
 Ile Asn Asp Asn Pro Pro Gln Ser Ser Gln Ser Ser Tyr Asp Val Tyr  
 450 455 460  
 Ile Glu Glu Asn Asn Leu Pro Gly Ala Pro Ile Leu Asn Leu Ser Val  
 465 470 475 480  
 Trp Asp Pro Asp Ala Pro Gln Asn Ala Arg Leu Ser Phe Phe Leu Leu  
 485 490 495  
 Glu Gln Gly Ala Glu Thr Gly Leu Val Gly Arg Tyr Phe Thr Ile Asn  
 500 505 510  
 Arg Asp Asn Gly Ile Val Ser Ser Leu Val Pro Leu Asp Tyr Glu Asp  
 515 520 525  
 Arg Arg Glu Phe Glu Leu Thr Ala His Ile Ser Asp Gly Gly Thr Pro  
 530 535 540  
 Val Leu Ala Thr Asn Ile Ser Val Asn Ile Phe Val Thr Asp Arg Asn  
 545 550 555 560  
 Asp Asn Ala Pro Gln Val Leu Tyr Pro Arg Pro Gly Gly Ser Ser Val  
 565 570 575  
 Glu Met Leu Pro Arg Gly Thr Ser Ala Gly His Leu Val Ser Arg Val  
 580 585 590  
 Val Gly Trp Asp Ala Asp Ala Gly His Asn Ala Trp Leu Ser Tyr Ser  
 595 600 605  
 Leu Phe Gly Ser Pro Asn Gln Ser Leu Phe Ala Ile Gly Leu His Thr  
 610 615 620  
 Gly Gln Ile Ser Thr Ala Arg Pro Val Gln Asp Thr Asp Ser Pro Arg  
 625 630 635 640  
 Gln Thr Leu Thr Val Leu Ile Lys Asp Asn Gly Glu Pro Ser Leu Ser  
 645 650 655  
 Thr Thr Ala Thr Leu Thr Val Ser Val Thr Glu Asp Ser Pro Glu Ala  
 660 665 670  
 Arg Ala Glu Phe Pro Ser Gly Ser Ala Pro Arg Glu Gln Lys Lys Asn  
 675 680 685  
 Leu Thr Phe Tyr Leu Leu Leu Ser Leu Ile Leu Val Ser Val Gly Phe  
 690 695 700

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Val Val Thr Val Phe Gly Val Ile Ile Phe Lys Val Tyr Lys Trp Lys  
705 710 715 720

Gln Ser Arg Asp Leu Tyr Arg Ala Pro Val Ser Ser Leu Tyr Arg Thr  
725 730 735

Pro Gly Pro Ser Leu His Ala Asp Ala Val Arg Gly Gly Leu Met Ser  
740 745 750

Pro His Leu Tyr His Gln Val Tyr Leu Thr Thr Asp Ser Arg Arg Ser  
755 760 765

Asp Pro Leu Leu Lys Lys Pro Gly Ala Ala Ser Pro Leu Ala Ser Arg  
770 775 780

Gln Asn Thr Leu Arg Ser Cys Asp Pro Val Phe Tyr Arg Gln Val Leu  
785 790 795 800

Gly Ala Glu Ser Ala Pro Pro Gly Gln Gln Ala Pro Pro Asn Thr Asp  
805 810 815

Trp Arg Phe Ser Gln Ala Gln Arg Pro Gly Thr Ser Gly Ser Gln Asn  
820 825 830

Gly Asp Asp Thr Gly Thr Trp Pro Asn Asn Gln Phe Asp Thr Glu Met  
835 840 845

Leu Gln Ala Met Ile Leu Ala Ser Ala Ser Glu Ala Ala Asp Gly Ser  
850 855 860

Ser Thr Leu Gly Gly Gly Ala Gly Thr Met Gly Leu Ser Ala Arg Tyr  
865 870 875 880

Gly Pro Gln Phe Thr Leu Gln His Val Pro Asp Tyr Arg Gln Asn Val  
885 890 895

Tyr Ile Pro Gly Ser Asn Ala His  
900

(2) INFORMATION FOR SEQ ID NO:98:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 441 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:98:

Asp Trp Val Ile Pro Pro Ile Asn Leu Pro Glu Asn Ser Arg Gly Pro  
1 5 10 15

Phe Pro Gln Glu Leu Val Arg Ile Arg Ser Asp Arg Asp Lys Asn Leu  
20 25 30

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Ser Leu Arg Tyr Thr Val Thr Gly Pro Gly Ala Asp Gln Pro Pro Thr  
35 40 45

Gly Ile Phe Ile Ile Asn Pro Ile Ser Gly Gln Leu Ser Val Thr Lys  
50 55 60

Pro Leu Asp Arg Glu Gln Ile Ala Arg Phe His Leu Arg Ala His Ala  
65 70 75 80

Val Asp Ile Asn Gly Asn Gln Val Glu Asn Pro Ile Asp Ile Val Ile  
85 90 95

Asn Val Ile Asp Met Asn Asp Asn Arg Pro Glu Phe Thr Ala Met Thr  
100 105 110

Phe Tyr Gly Glu Val Pro Glu Asn Arg Val Asp Ile Ile Val Ala Asn  
115 120 125

Leu Thr Val Thr Asp Lys Asp Gln Pro His Thr Pro Ala Trp Asn Ala  
130 135 140

Val Thr Arg Ile Ser Gly Gly Asp Pro Thr Gly Arg Phe Ala Ile Gln  
145 150 155 160

Thr Asp Pro Asn Ser Asn Asp Gly Leu Val Thr Val Val Lys Pro Ile  
165 170 175

Asp Phe Glu Thr Asn Arg Met Phe Val Leu Thr Val Ala Ala Glu Asn  
180 185 190

Gln Val Pro Leu Ala Lys Gly Ile Gln His Pro Pro Gln Ser Thr Ala  
195 200 205

Thr Val Ser Val Thr Val Ile Asp Val Asn Glu Asn Pro Tyr Phe Ala  
210 215 220

Pro Asn Pro Lys Ile Ile Arg Gln Glu Glu Gly Leu His Ala Gly Thr  
225 230 235 240

Met Leu Thr Thr Phe Thr Ala Gly Asp Pro Asp Arg Tyr Met Gln Gln  
245 250 255

Asn Ile Arg Tyr Thr Lys Leu Ser Asp Pro Ala Asn Trp Leu Lys Ile  
260 265 270

Asp Pro Val Asn Gly Gln Ile Thr Thr Ile Ala Val Leu Asp Arg Glu  
275 280 285

Ser Pro Asn Val Lys Asn Asn Ile Tyr Asn Ala Thr Phe Leu Ala Ser  
290 295 300

Asp Asn Gly Ile Pro Pro Met Ser Gly Thr Gly Thr Leu Gln Ile Tyr  
305 310 315 320

Leu Leu Asp Ile Asn Asp Asn Ala Pro Gln Val Leu Pro Gln Glu Ala  
325 330 335

Glu Thr Cys Glu Thr Pro Asp Pro Asn Ser Ile Asn Ile Thr Thr Ala  
340 345 350

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Leu Asp Tyr Asp Ile Asp Pro Asn Ala Gly Pro Phe Ala Tyr Asp Leu  
 355 360 365  
 Pro Leu Ser Pro Val Thr Ile Lys Arg Asn Trp Thr Ile Thr Arg Leu  
 370 375 380  
 Asn Gly Asp Phe Ala Gln Leu Asn Leu Lys Ile Lys Phe Leu Glu Ala  
 385 390 395 400  
 Gly Ile Tyr Glu Val Pro Ile Ile Ile Thr Asp Ser Gly Asn Pro Pro  
 405 410 415  
 Lys Ser Asn Lys Ser Ile Leu Arg Val Arg Val Cys Gln Cys Asp Phe  
 420 425 430  
 Asn Gly Asp Cys Thr Asp Val Asp Arg  
 435 440

(2) INFORMATION FOR SEQ ID NO:99:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 105 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:99:

Glu Asp Thr Val Tyr Ser Phe Asp Ile Pro Glu Asn Ala Gln Arg Gly  
 1 5 10 15  
 Tyr Gln Val Gly Gln Ile Val Ala Arg Asp Ala Asp Leu Gly Gln Asn  
 20 25 30  
 Ala Gln Leu Ser Tyr Gly Val Val Ser Asp Trp Ala Asn Asp Val Phe  
 35 40 45  
 Ser Leu Asn Pro Gln Thr Gly Met Leu Thr Leu Thr Ala Arg Leu Asp  
 50 55 60  
 Tyr Glu Glu Val Gln His Tyr Ile Leu Ile Val Gln Ala Gln Asp Asn  
 65 70 75 80  
 Gly Gln Pro Ser Leu Ser Thr Thr Ile Thr Val Tyr Cys Asn Val Leu  
 85 90 95  
 Asp Leu Asn Asp Asn Ala Pro Ile Phe  
 100 105

(2) INFORMATION FOR SEQ ID NO:100:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 7 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

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(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:100:

Asp Xaa Asp Xaa Gly Xaa Asn  
1 5

(2) INFORMATION FOR SEQ ID NO:101:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 7 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:101:

Ala Xaa Asp Xaa Gly Xaa Pro  
1 5

(2) INFORMATION FOR SEQ ID NO:102:

(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 4650 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

(A) NAME/KEY: CDS  
(B) LOCATION: 495..4103

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:102:

CCTCTATTCG ACATTCTCTT TGGATTGTTT TGCTATAACT TGAAATTTGG GATGTCACAA	60
ACGAAACTGT CATCTGTTTC CGCCAAACTG TGGTTCTGCT AATCTCCCAG GCTGGCAGCA	120
TTGGAGACTT GCTGACTTCT TTCATCCCCC ACTCTTTTCA CCTGAAATTC CTTTCCTTGG	180
TTTTGCTCTA AGTCCTATGC TTCAGTCAGG GGCCAACCAA ATCTCACTGC CTCCTTTTTA	240
TCATGAAGCC TTTGATCACT GATAGTTCTT TTTATATCTT GAAAAATCAC CCTTCCCAGT	300
ACAGTTAATA TTTAGTATCT CTA CTACTCATCT TGGCACTTAC TCACAGCTCC ATAATTCACT	360
CGTTTTTCGTA CCTCTTCATG GTGATGGGGA GCCCTTTGGA GGTGGTGA CTGTCTTTATA	420
CTCCTCATGA TGCTTCACAT GTGGCAGGCG TGGAGTGCCC GGAGGCGGCC CTCCTGATTC	480

TGGGGCCTCC	CAGG	ATG	GAG	CCC	CTG	AGG	CAC	AGC	CCA	GGC	CCT	GGG	GGG	530
	Met	Glu	Pro	Leu	Arg	His	Ser	Pro	Gly	Pro	Gly	Gly		
	1				5					10				
CAA CGG CTA CTG CTG CCC TCC ATG CTG CTA GCA CTG CTG CTC CTG CTG														578
Gln Arg Leu Leu Leu Pro Ser Met Leu Leu Ala Leu Leu Leu Leu														
	15					20				25				
GCT CCA TCC CCA GGC CAC GCC ACT CGG GTA GTG TAC AAG GTG CCG GAG														626
Ala Pro Ser Pro Gly His Ala Thr Arg Val Val Tyr Lys Val Pro Glu														
	30					35				40				
GAA CAG CCA CCC AAC ACC CTC ATT GGG AGC CTC GCA GCC GAC TAT GGT														674
Glu Gln Pro Pro Asn Thr Leu Ile Gly Ser Leu Ala Ala Asp Tyr Gly														
	45				50				55				60	
TTT CCA GAT GTG GGG CAC CTG TAC AAG CTA GAG GTG GGT GCC CCG TAC														722
Phe Pro Asp Val Gly His Leu Tyr Lys Leu Glu Val Gly Ala Pro Tyr														
				65				70					75	
CTT CGC GTG GAT GGC AAG ACA GGT GAC ATT TTC ACC ACC GAG ACC TCC														770
Leu Arg Val Asp Gly Lys Thr Gly Asp Ile Phe Thr Thr Glu Thr Ser														
			80				85					90		
ATC GAC CGT GAG GGG CTC CGT GAA TGC CAG AAC CAG CTC CCT GGT GAT														818
Ile Asp Arg Glu Gly Leu Arg Glu Cys Gln Asn Gln Leu Pro Gly Asp														
		95					100					105		
CCC TGC ATC CTG GAG TTT GAG GTA TCT ATC ACA GAC CTC GTG CAG AAT														866
Pro Cys Ile Leu Glu Phe Glu Val Ser Ile Thr Asp Leu Val Gln Asn														
	110					115				120				
GCG AGC CCC CGG CTG CTA GAG GGC CAG ATA GAA GTA CAA GAC ATC AAT														914
Ala Ser Pro Arg Leu Glu Gly Gln Ile Glu Val Gln Asp Ile Asn														
	125				130				135				140	
GAC AAC ACA CCC AAC TTC GCC TCA CCA GTC ATC ACT CTG GCC ATC CCT														962
Asp Asn Thr Pro Asn Phe Ala Ser Pro Val Ile Thr Leu Ala Ile Pro														
				145				150					155	
GAG AAC ACC AAC ATC GGC TCA CTC TTC CCC ATC CCG CTG GCT TCA GAC														1010
Glu Asn Thr Asn Ile Gly Ser Leu Phe Pro Ile Pro Leu Ala Ser Asp														
			160				165					170		
CGT GAT GCT GGT CCC AAC GGT GTG GCA TCC TAT GAG CTG CAG GTG GCA														1058
Arg Asp Ala Gly Pro Asn Gly Val Ala Ser Tyr Glu Leu Gln Val Ala														
		175				180					185			
GAG GAC CAG GAG GAG AAG CAA CCA CAG CTC ATT GTG ATG GGC AAC CTG														1106
Glu Asp Gln Glu Glu Lys Gln Pro Gln Leu Ile Val Met Gly Asn Leu														
	190					195				200				
GAC CGT GAG CGC TGG GAC TCC TAT GAC CTC ACC ATC AAG GTG CAG GAT														1154
Asp Arg Glu Arg Trp Asp Ser Tyr Asp Leu Thr Ile Lys Val Gln Asp														
	205				210				215				220	
GGC GGC AGC CCC CCA CGC GCC ACG AGT GCC CTG CTG CGT GTC ACC GTG														1202
Gly Gly Ser Pro Pro Arg Ala Thr Ser Ala Leu Leu Arg Val Thr Val														
			225					230					235	

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CTT GAC ACC AAT GAC AAC GCC CCC AAG TTT GAG CGG CCC TCC TAT GAG Leu Asp Thr Asn Asp Asn Ala Pro Lys Phe Glu Arg Pro Ser Tyr Glu 240 245 250	1250
GCC GAA CTA TCT GAG AAT AGC CCC ATA GGC CAC TCG GTC ATC CAG GTG Ala Glu Leu Ser Glu Asn Ser Pro Ile Gly His Ser Val Ile Gln Val 255 260 265	1298
AAG GCC AAT GAC TCA GAC CAA GGT GCC AAT GCA GAA ATC GAA TAC ACA Lys Ala Asn Asp Ser Asp Gln Gly Ala Asn Ala Glu Ile Glu Tyr Thr 270 275 280	1346
TTC CAC CAG GCG CCC GAA GTT GTG AGG CGT CTT CTT CGA CTG GAC AGG Phe His Gln Ala Pro Glu Val Val Arg Arg Leu Leu Arg Leu Asp Arg 285 290 295 300	1394
AAC ACT GGA CTT ATC ACT GTT CAG GGC CCG GTG GAC CGT GAG GAC CTA Asn Thr Gly Leu Ile Thr Val Gln Gly Pro Val Asp Arg Glu Asp Leu 305 310 315	1442
AGC ACC CTG CGC TTC TCA GTG CTT GCT AAG GAC CGA GGC ACC AAC CCC Ser Thr Leu Arg Phe Ser Val Leu Ala Lys Asp Arg Gly Thr Asn Pro 320 325 330	1490
AAG AGT GCC CGT GCC CAG GTG GTT GTG ACC GTG AAG GAC ATG AAT GAC Lys Ser Ala Arg Ala Gln Val Val Val Thr Val Lys Asp Met Asn Asp 335 340 345	1538
AAT GCC CCC ACC ATT GAG ATC CGG GGC ATA GGG CTA GTG ACT CAT CAA Asn Ala Pro Thr Ile Glu Ile Arg Gly Ile Gly Leu Val Thr His Gln 350 355 360	1586
GAT GGG ATG GCT AAC ATC TCA GAG GAT GTG GCA GAG GAG ACA GCT GTG Asp Gly Met Ala Asn Ile Ser Glu Asp Val Ala Glu Glu Thr Ala Val 365 370 375 380	1634
GCC CTG GTG CAG GTG TCT GAC CGA GAT GAG GGA GAG AAT GCA GCT GTC Ala Leu Val Gln Val Ser Asp Arg Asp Glu Gly Glu Asn Ala Ala Val 385 390 395	1682
ACC TGT GTG GTG GCA GGT GAT GTG CCC TTC CAG CTG CGC CAG GCC AGT Thr Cys Val Val Ala Gly Asp Val Pro Phe Gln Leu Arg Gln Ala Ser 400 405 410	1730
GAG ACA GGC AGT GAC AGC AAG AAG AAG TAT TTC CTG CAG ACT ACC ACC Glu Thr Gly Ser Asp Ser Lys Lys Lys Tyr Phe Leu Gln Thr Thr Thr 415 420 425	1778
CCG CTA GAC TAC GAG AAG GTC AAA GAC TAC ACC ATT GAG ATT GTG GCT Pro Leu Asp Tyr Glu Lys Val Lys Asp Tyr Thr Ile Glu Ile Val Ala 430 435 440	1826
GTG GAC TCT GGC AAC CCC CCA CTC TCC AGC ACT AAC TCC CTC AAG GTG Val Asp Ser Gly Asn Pro Pro Leu Ser Ser Thr Asn Ser Leu Lys Val 445 450 455 460	1874
CAG GTG GTG GAC GTC AAT GAC AAC GCA CCT GTC TTC ACT CAG AGT GTC Gln Val Val Asp Val Asn Asp Asn Ala Pro Val Phe Thr Gln Ser Val 465 470 475	1922

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ACT	GAG	GTC	GCC	TTC	CCG	GAA	AAC	AAC	AAG	CCT	GGT	GAA	GTG	ATT	GCT	1970
Thr	Glu	Val	Ala	Phe	Pro	Glu	Asn	Asn	Lys	Pro	Gly	Glu	Val	Ile	Ala	
			480					485					490			
GAG	ATC	ACT	GCC	AGT	GAT	GCT	GAC	TCT	GGC	TCT	AAT	GCT	GAG	CTG	GTT	2018
Glu	Ile	Thr	Ala	Ser	Asp	Ala	Asp	Ser	Gly	Ser	Asn	Ala	Glu	Leu	Val	
		495				500						505				
TAC	TCT	CTG	GAG	CCT	GAG	CCG	GCT	GCT	AAG	GGC	CTC	TTC	ACC	ATC	TCA	2066
Tyr	Ser	Leu	Glu	Pro	Glu	Pro	Ala	Ala	Lys	Gly	Leu	Phe	Thr	Ile	Ser	
	510					515					520					
CCC	GAG	ACT	GGA	GAG	ATC	CAG	GTG	AAG	ACA	TCT	CTG	GAT	CGG	GAA	CAG	2114
Pro	Glu	Thr	Gly	Glu	Ile	Gln	Val	Lys	Thr	Ser	Leu	Asp	Arg	Glu	Gln	
525					530					535					540	
CGG	GAG	AGC	TAT	GAG	TTG	AAG	GTG	GTG	GCA	GCT	GAC	CGG	GGC	AGT	CCT	2162
Arg	Glu	Ser	Tyr	Glu	Leu	Lys	Val	Val	Ala	Ala	Asp	Arg	Gly	Ser	Pro	
				545					550					555		
AGC	CTC	CAG	GGC	ACA	GCC	ACT	GTC	CTT	GTC	AAT	GTG	CTG	GAC	TGC	AAT	2210
Ser	Leu	Gln	Gly	Thr	Ala	Thr	Val	Leu	Val	Asn	Val	Leu	Asp	Cys	Asn	
			560					565					570			
GAC	AAT	GAC	CCC	AAA	TTT	ATG	CTG	AGT	GGC	TAC	AAC	TTC	TCA	GTG	ATG	2258
Asp	Asn	Asp	Pro	Lys	Phe	Met	Leu	Ser	Gly	Tyr	Asn	Phe	Ser	Val	Met	
		575					580					585				
GAG	AAC	ATG	CCA	GCA	CTG	AGT	CCA	GTG	GGC	ATG	GTG	ACT	GTC	ATT	GAT	2306
Glu	Asn	Met	Pro	Ala	Leu	Ser	Pro	Val	Gly	Met	Val	Thr	Val	Ile	Asp	
	590					595					600					
GGA	GAC	AAG	GGG	GAG	AAT	GCC	CAG	GTG	CAG	CTC	TCA	GTG	GAG	CAG	GAC	2354
Gly	Asp	Lys	Gly	Glu	Asn	Ala	Gln	Val	Gln	Leu	Ser	Val	Glu	Gln	Asp	
605					610					615					620	
AAC	GGT	GAC	TTT	GTT	ATC	CAG	AAT	GGC	ACA	GGC	ACC	ATC	CTA	TCC	AGC	2402
Asn	Gly	Asp	Phe	Val	Ile	Gln	Asn	Gly	Thr	Gly	Thr	Ile	Leu	Ser	Ser	
				625					630					635		
CTG	AGC	TTT	GAT	CGA	GAG	CAA	CAA	AGC	ACC	TAC	ACC	TTC	CAG	CTG	AAG	2450
Leu	Ser	Phe	Asp	Arg	Glu	Gln	Gln	Ser	Thr	Tyr	Thr	Phe	Gln	Leu	Lys	
			640					645					650			
GCA	GTG	GAT	GGT	GGC	GTC	CCA	CCT	CGC	TCA	GCT	TAC	GTT	GGT	GTC	ACC	2498
Ala	Val	Asp	Gly	Gly	Val	Pro	Pro	Arg	Ser	Ala	Tyr	Val	Gly	Val	Thr	
		655				660						665				
ATC	AAT	GTG	CTG	GAC	GAG	AAT	GAC	AAC	GCA	CCC	TAT	ATC	ACT	GCC	CCT	2546
Ile	Asn	Val	Leu	Asp	Glu	Asn	Asp	Asn	Ala	Pro	Tyr	Ile	Thr	Ala	Pro	
	670					675					680					
TCT	AAC	ACC	TCT	CAC	AAG	CTG	CTG	ACC	CCC	CAG	ACA	CGT	CTT	GGT	GAG	2594
Ser	Asn	Thr	Ser	His	Lys	Leu	Leu	Thr	Pro	Gln	Thr	Arg	Leu	Gly	Glu	
685					690					695					700	
ACG	GTC	AGC	CAG	GTG	GCA	GCC	GAG	GAC	TTT	GAC	TCT	GGT	GTC	AAT	GCC	2642
Thr	Val	Ser	Gln	Val	Ala	Ala	Glu	Asp	Phe	Asp	Ser	Gly	Val	Asn	Ala	
				705					710					715		

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GAG CTG ATC TAC AGC ATT GCA GGT GGC AAC CCT TAT GGA CTC TTC CAG Glu Leu Ile Tyr Ser Ile Ala Gly Gly Asn Pro Tyr Gly Leu Phe Gln 720 725 730	2690
ATT GGG TCA CAT TCA GGT GCC ATC ACC CTG GAG AAG GAG ATT GAG CGG Ile Gly Ser His Ser Gly Ala Ile Thr Leu Glu Lys Glu Ile Glu Arg 735 740 745	2738
CGC CAC CAT GGG CTA CAC CGC CTG GTG GTG AAG GTC AGT GAC CGC GGC Arg His His Gly Leu His Arg Leu Val Val Lys Val Ser Asp Arg Gly 750 755 760	2786
AAG CCC CCA CGC TAT GGC ACA GCC TTG GTC CAT CTT TAT GTC AAT GAG Lys Pro Pro Arg Tyr Gly Thr Ala Leu Val His Leu Tyr Val Asn Glu 765 770 775 780	2834
ACT CTG GCC AAC CGC ACG CTG CTG GAG ACC CTC CTG GGC CAC AGC CTG Thr Leu Ala Asn Arg Thr Leu Leu Glu Thr Leu Leu Gly His Ser Leu 785 790 795	2882
GAC ACG CCG CTG GAT ATT GAC ATT GCT GGG GAT CCA GAA TAT GAG CGC Asp Thr Pro Leu Asp Ile Asp Ile Ala Gly Asp Pro Glu Tyr Glu Arg 800 805 810	2930
TCC AAG CAG CGT GGC AAC ATT CTC TTT GGT GTG GTG GCT GGT GTG GTG Ser Lys Gln Arg Gly Asn Ile Leu Phe Gly Val Val Ala Gly Val Val 815 820 825	2978
GCC GTG GCC TTG CTC ATC GCC CTG GCG GTT CTT GTG CGC TAC TGC AGA Ala Val Ala Leu Leu Ile Ala Leu Ala Val Leu Val Arg Tyr Cys Arg 830 835 840	3026
CAG CGG GAG GCC AAA AGT GGT TAC CAG GCT GGT AAG AAG GAG ACC AAG Gln Arg Glu Ala Lys Ser Gly Tyr Gln Ala Gly Lys Lys Glu Thr Lys 845 850 855 860	3074
GAC CTG TAT GCC CCC AAG CCC AGT GGC AAG GCC TCC AAG GGA AAC AAA Asp Leu Tyr Ala Pro Lys Pro Ser Gly Lys Ala Ser Lys Gly Asn Lys 865 870 875	3122
AGC AAA GGC AAG AAG AGC AAG TCC CCA AAG CCC GTG AAG CCA GTG GAG Ser Lys Gly Lys Lys Ser Lys Ser Pro Lys Pro Val Lys Pro Val Glu 880 885 890	3170
GAC GAG GAT GAG GCC GGG CTG CAG AAG TCC CTC AAG TTC AAC CTG ATG Asp Glu Asp Glu Ala Gly Leu Gln Lys Ser Leu Lys Phe Asn Leu Met 895 900 905	3218
AGC GAT GCC CCT GGG GAC AGT CCC CGC ATC CAC CTG CCC CTC AAC TAC Ser Asp Ala Pro Gly Asp Ser Pro Arg Ile His Leu Pro Leu Asn Tyr 910 915 920	3266
CCA CCA GGC AGC CCT GAC CTG GGC CGC CAC TAT CGC TCT AAC TCC CCA Pro Pro Gly Ser Pro Asp Leu Gly Arg His Tyr Arg Ser Asn Ser Pro 925 930 935 940	3314
CTG CCT TCC ATC CAG CTG CAG CCC CAG TCA CCC TCA GCC TCC AAG AAG Leu Pro Ser Ile Gln Leu Gln Pro Gln Ser Pro Ser Ala Ser Lys Lys 945 950 955	3362

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CAC	CAG	GTG	GTA	CAG	GAC	CTG	CCA	CCT	GCA	AAC	ACA	TTC	GTG	GGC	ACC	3410
His	Gln	Val	Val	Gln	Asp	Leu	Pro	Pro	Ala	Asn	Thr	Phe	Val	Gly	Thr	
			960					965					970			
GGG	GAC	ACC	ACG	TCC	ACG	GGC	TCT	GAG	CAG	TAC	TCC	GAC	TAC	AGC	TAC	3458
Gly	Asp	Thr	Thr	Ser	Thr	Gly	Ser	Glu	Gln	Tyr	Ser	Asp	Tyr	Ser	Tyr	
		975					980					985				
CGC	ACC	AAC	CCC	CCC	AAA	TAC	CCC	AGC	AAG	CAG	TTA	CCT	CAC	CGC	CGC	3506
Arg	Thr	Asn	Pro	Pro	Lys	Tyr	Pro	Ser	Lys	Gln	Leu	Pro	His	Arg	Arg	
	990					995					1000					
GTC	ACC	TTC	TCG	GCC	ACC	AGC	CAG	GCC	CAG	GAG	CTG	CAG	GAC	CCA	TCC	3554
Val	Thr	Phe	Ser	Ala	Thr	Ser	Gln	Ala	Gln	Glu	Leu	Gln	Asp	Pro	Ser	
1005					1010					1015					1020	
CAG	CAC	AGT	TAC	TAT	GAC	AGT	GGC	CTG	GAG	GAG	TCT	GAG	ACG	CCG	TCC	3602
Gln	His	Ser	Tyr	Tyr	Asp	Ser	Gly	Leu	Glu	Glu	Ser	Glu	Thr	Pro	Ser	
				1025					1030					1035		
AGC	AAG	TCA	TCC	TCA	GGG	CCT	CGA	CTC	GGT	CCC	CTG	GCC	CTG	CCT	GAG	3650
Ser	Lys	Ser	Ser	Ser	Gly	Pro	Arg	Leu	Gly	Pro	Leu	Ala	Leu	Pro	Glu	
			1040					1045					1050			
GAT	CAC	TAT	GAG	CGC	ACC	ACC	CCT	GAT	GGC	AGC	ATA	GGA	GAG	ATG	GAG	3698
Asp	His	Tyr	Glu	Arg	Thr	Thr	Pro	Asp	Gly	Ser	Ile	Gly	Glu	Met	Glu	
		1055					1060					1065				
CAC	CCC	GAG	AAT	GAC	CTT	CGC	CCT	TTG	CCT	GAT	GTC	GCC	ATG	ACA	CGC	3746
His	Pro	Glu	Asn	Asp	Leu	Arg	Pro	Leu	Pro	Asp	Val	Ala	Met	Thr	Gly	
	1070					1075					1080					
ACA	TGT	ACC	CGG	GAG	TGC	AGT	GAG	TTT	GGC	CAC	TCT	GAC	ACA	TGC	TGG	3794
Thr	Cys	Thr	Arg	Glu	Cys	Ser	Glu	Phe	Gly	His	Ser	Asp	Thr	Cys	Trp	
1085					1090					1095					1100	
ATG	CCT	GGC	CAG	TCA	TCT	CCC	AGC	CGC	CGG	ACC	AAG	AGC	AGC	GCC	CTC	3842
Met	Pro	Gly	Gln	Ser	Ser	Pro	Ser	Arg	Arg	Thr	Lys	Ser	Ser	Ala	Leu	
				1105					1110					1115		
AAA	CTC	TCC	ACC	TTC	ATG	CCT	TAC	CAG	GAC	CGA	GGA	GGG	CAG	GAG	CCT	3890
Lys	Leu	Ser	Thr	Phe	Met	Pro	Tyr	Gln	Asp	Arg	Gly	Gly	Gln	Glu	Pro	
			1120					1125					1130			
GCG	GGC	GCC	GGC	AGC	CCC	AGC	CCC	CCG	GAA	GAC	CGG	AAC	ACC	AAA	ACG	3938
Ala	Gly	Ala	Gly	Ser	Pro	Ser	Pro	Pro	Glu	Asp	Arg	Asn	Thr	Lys	Thr	
		1135					1140					1145				
GCC	CCC	GTG	CGC	CTC	CTG	CCC	TCC	TAC	AGT	GCC	TTC	TCC	CAC	AGT	AGC	3986
Ala	Pro	Val	Arg	Leu	Leu	Pro	Ser	Tyr	Ser	Ala	Phe	Ser	His	Ser	Ser	
	1150					1155					1160					
CAT	GAT	TCC	TGC	AAG	GAC	TCG	GCC	ACC	TTG	GAG	GAA	ATC	CCC	CTG	ACC	4034
His	Asp	Ser	Cys	Lys	Asp	Ser	Ala	Thr	Leu	Glu	Glu	Ile	Pro	Leu	Thr	
1165					1170					1175				1180		
CAG	ACC	TCG	GAC	TTC	CCA	CCC	GCA	GCC	ACA	CCG	GCA	TCT	GCC	CAG	ACG	4082
Gln	Thr	Ser	Asp	Phe	Pro	Pro	Ala	Ala	Thr	Pro	Ala	Ser	Ala	Gln	Thr	
				1185					1190					1195		

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GCC AAG CGC GAG ATC TAC CTG TGAGCCCCCT ACTGGCCGGC CCCCCTCCCC 4133  
Ala Lys Arg Glu Ile Tyr Leu  
1200

CAGCGCCGGC CAGCTCCCAA ATGCCCCATTC CAGGGCCTCA CTCTCCACCC CTTCAGCGTG 4193  
GACTTCCTGC CAGGGCCCAA GTGGGGGTAT CACTGACCTC ATGACCACGC TGGCCCTTCT 4253  
CCCATGCAGG GTCCAGGTCC TCTCCCCTCA TTTCCATCTC CCAGCCCAGG GGCCCTTCC 4313  
CCTTTATGGG GCTTCCCCCA GCTGATGCCC AAGAGGGGCTC CTCTGCAATG ACTGGGCTCC 4373  
TTCCCTTGAC TTCCAGGGAG CACCCCCTCG ATTTGGGCAG ATGGTGGAGT CAAGGGTGGG 4433  
CAGCGTACTT CTAATCATT GTTTCCTCA TGGCCGACCA GGGCGGGGAT AGCATGCCCA 4493  
ATTTTAGCCC TGAAGCAGGG CTGAAGTGGG GAGCCCCCTT CCCTGGGAGC TCCCAGAGGA 4553  
AACTCTTGAC CACCAGTGGC TCCCTGAAGG GCTTTTGTTA CCAAAGGTGG GGTAGGGACG 4613  
GGGGTGGGAG TGGAGCGGAG GCCTTGTTTT CCCGTGG 4650

(2) INFORMATION FOR SEQ ID NO:103:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 1203 amino acids  
(B) TYPE: amino acid  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:103:

Met Glu Pro Leu Arg His Ser Pro Gly Pro Gly Gly Gln Arg Leu Leu  
1 5 10 15  
Leu Pro Ser Met Leu Leu Ala Leu Leu Leu Leu Ala Pro Ser Pro  
20 25 30  
Gly His Ala Thr Arg Val Val Tyr Lys Val Pro Glu Glu Gln Pro Pro  
35 40 45  
Asn Thr Leu Ile Gly Ser Leu Ala Ala Asp Tyr Gly Phe Pro Asp Val  
50 55 60  
Gly His Leu Tyr Lys Leu Glu Val Gly Ala Pro Tyr Leu Arg Val Asp  
65 70 75 80  
Gly Lys Thr Gly Asp Ile Phe Thr Thr Glu Thr Ser Ile Asp Arg Glu  
85 90 95  
Gly Leu Arg Glu Cys Gln Asn Gln Leu Pro Gly Asp Pro Cys Ile Leu  
100 105 110  
Glu Phe Glu Val Ser Ile Thr Asp Leu Val Gln Asn Ala Ser Pro Arg  
115 120 125



Leu Leu Glu Gly Gln Ile Glu Val Gln Asp Ile Asn Asp Asn Thr Pro  
 130 135 140  
 Asn Phe Ala Ser Pro Val Ile Thr Leu Ala Ile Pro Glu Asn Thr Asn  
 145 150 155 160  
 Ile Gly Ser Leu Phe Pro Ile Pro Leu Ala Ser Asp Arg Asp Ala Gly  
 165 170 175  
 Pro Asn Gly Val Ala Ser Tyr Glu Leu Gln Val Ala Glu Asp Gln Glu  
 180 185 190  
 Glu Lys Gln Pro Gln Leu Ile Val Met Gly Asn Leu Asp Arg Glu Arg  
 195 200 205  
 Trp Asp Ser Tyr Asp Leu Thr Ile Lys Val Gln Asp Gly Gly Ser Pro  
 210 215 220  
 Pro Arg Ala Thr Ser Ala Leu Leu Arg Val Thr Val Leu Asp Thr Asn  
 225 230 235 240  
 Asp Asn Ala Pro Lys Phe Glu Arg Pro Ser Tyr Glu Ala Glu Leu Ser  
 245 250 255  
 Glu Asn Ser Pro Ile Gly His Ser Val Ile Gln Val Lys Ala Asn Asp  
 260 265 270  
 Ser Asp Gln Gly Ala Asn Ala Glu Ile Glu Tyr Thr Phe His Gln Ala  
 275 280 285  
 Pro Glu Val Val Arg Arg Leu Leu Arg Leu Asp Arg Asn Thr Gly Leu  
 290 295 300  
 Ile Thr Val Gln Gly Pro Val Asp Arg Glu Asp Leu Ser Thr Leu Arg  
 305 310 315 320  
 Phe Ser Val Leu Ala Lys Asp Arg Gly Thr Asn Pro Lys Ser Ala Arg  
 325 330 335  
 Ala Gln Val Val Val Thr Val Lys Asp Met Asn Asp Asn Ala Pro Thr  
 340 345 350  
 Ile Glu Ile Arg Gly Ile Gly Leu Val Thr His Gln Asp Gly Met Ala  
 355 360 365  
 Asn Ile Ser Glu Asp Val Ala Glu Glu Thr Ala Val Ala Leu Val Gln  
 370 375 380  
 Val Ser Asp Arg Asp Glu Gly Glu Asn Ala Ala Val Thr Cys Val Val  
 385 390 395 400  
 Ala Gly Asp Val Pro Phe Gln Leu Arg Gln Ala Ser Glu Thr Gly Ser  
 405 410 415  
 Asp Ser Lys Lys Lys Tyr Phe Leu Gln Thr Thr Thr Pro Leu Asp Tyr  
 420 425 430  
 Glu Lys Val Lys Asp Tyr Thr Ile Glu Ile Val Ala Val Asp Ser Gly  
 435 440 445

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Asn Pro Pro Leu Ser Ser Thr Asn Ser Leu Lys Val Gln Val Val Asp  
 450 455 460  
 Val Asn Asp Asn Ala Pro Val Phe Thr Gln Ser Val Thr Glu Val Ala  
 465 470 475 480  
 Phe Pro Glu Asn Asn Lys Pro Gly Glu Val Ile Ala Glu Ile Thr Ala  
 485 490 495  
 Ser Asp Ala Asp Ser Gly Ser Asn Ala Glu Leu Val Tyr Ser Leu Glu  
 500 505 510  
 Pro Glu Pro Ala Ala Lys Gly Leu Phe Thr Ile Ser Pro Glu Thr Gly  
 515 520 525  
 Glu Ile Gln Val Lys Thr Ser Leu Asp Arg Glu Gln Arg Glu Ser Tyr  
 530 535 540  
 Glu Leu Lys Val Val Ala Ala Asp Arg Gly Ser Pro Ser Leu Gln Gly  
 545 550 555 560  
 Thr Ala Thr Val Leu Val Asn Val Leu Asp Cys Asn Asp Asn Asp Pro  
 565 570 575  
 Lys Phe Met Leu Ser Gly Tyr Asn Phe Ser Val Met Glu Asn Met Pro  
 580 585 590  
 Ala Leu Ser Pro Val Gly Met Val Thr Val Ile Asp Gly Asp Lys Gly  
 595 600 605  
 Glu Asn Ala Gln Val Gln Leu Ser Val Glu Gln Asp Asn Gly Asp Phe  
 610 615 620  
 Val Ile Gln Asn Gly Thr Gly Thr Ile Leu Ser Ser Leu Ser Phe Asp  
 625 630 635 640  
 Arg Glu Gln Gln Ser Thr Tyr Thr Phe Gln Leu Lys Ala Val Asp Gly  
 645 650 655  
 Gly Val Pro Pro Arg Ser Ala Tyr Val Gly Val Thr Ile Asn Val Leu  
 660 665 670  
 Asp Glu Asn Asp Asn Ala Pro Tyr Ile Thr Ala Pro Ser Asn Thr Ser  
 675 680 685  
 His Lys Leu Leu Thr Pro Gln Thr Arg Leu Gly Glu Thr Val Ser Gln  
 690 695 700  
 Val Ala Ala Glu Asp Phe Asp Ser Gly Val Asn Ala Glu Leu Ile Tyr  
 705 710 715 720  
 Ser Ile Ala Gly Gly Asn Pro Tyr Gly Leu Phe Gln Ile Gly Ser His  
 725 730 735  
 Ser Gly Ala Ile Thr Leu Glu Lys Glu Ile Glu Arg Arg His His Gly  
 740 745 750  
 Leu His Arg Leu Val Val Lys Val Ser Asp Arg Gly Lys Pro Pro Arg  
 755 760 765

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Tyr Gly Thr Ala Leu Val His Leu Tyr Val Asn Glu Thr Leu Ala Asn  
 770 775 780  
 Arg Thr Leu Leu Glu Thr Leu Leu Gly His Ser Leu Asp Thr Pro Leu  
 785 790 795 800  
 Asp Ile Asp Ile Ala Gly Asp Pro Glu Tyr Glu Arg Ser Lys Gln Arg  
 805 810 815  
 Gly Asn Ile Leu Phe Gly Val Val Ala Gly Val Val Ala Val Ala Leu  
 820 825 830  
 Leu Ile Ala Leu Ala Val Leu Val Arg Tyr Cys Arg Gln Arg Glu Ala  
 835 840 845  
 Lys Ser Gly Tyr Gln Ala Gly Lys Lys Glu Thr Lys Asp Leu Tyr Ala  
 850 855 860  
 Pro Lys Pro Ser Gly Lys Ala Ser Lys Gly Asn Lys Ser Lys Gly Lys  
 865 870 875 880  
 Lys Ser Lys Ser Pro Lys Pro Val Lys Pro Val Glu Asp Glu Asp Glu  
 885 890 895  
 Ala Gly Leu Gln Lys Ser Leu Lys Phe Asn Leu Met Ser Asp Ala Pro  
 900 905 910  
 Gly Asp Ser Pro Arg Ile His Leu Pro Leu Asn Tyr Pro Pro Gly Ser  
 915 920 925  
 Pro Asp Leu Gly Arg His Tyr Arg Ser Asn Ser Pro Leu Pro Ser Ile  
 930 935 940  
 Gln Leu Gln Pro Gln Ser Pro Ser Ala Ser Lys Lys His Gln Val Val  
 945 950 955 960  
 Gln Asp Leu Pro Pro Ala Asn Thr Phe Val Gly Thr Gly Asp Thr Thr  
 965 970 975  
 Ser Thr Gly Ser Glu Gln Tyr Ser Asp Tyr Ser Tyr Arg Thr Asn Pro  
 980 985 990  
 Pro Lys Tyr Pro Ser Lys Gln Leu Pro His Arg Arg Val Thr Phe Ser  
 995 1000 1005  
 Ala Thr Ser Gln Ala Gln Glu Leu Gln Asp Pro Ser Gln His Ser Tyr  
 1010 1015 1020  
 Tyr Asp Ser Gly Leu Glu Glu Ser Glu Thr Pro Ser Ser Lys Ser Ser  
 1025 1030 1035 1040  
 Ser Gly Pro Arg Leu Gly Pro Leu Ala Leu Pro Glu Asp His Tyr Glu  
 1045 1050 1055  
 Arg Thr Thr Pro Asp Gly Ser Ile Gly Glu Met Glu His Pro Glu Asn  
 1060 1065 1070  
 Asp Leu Arg Pro Leu Pro Asp Val Ala Met Thr Gly Thr Cys Thr Arg  
 1075 1080 1085

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Glu Cys Ser Glu Phe Gly His Ser Asp Thr Cys Trp Met Pro Gly Gln  
 1090 1095 1100

Ser Ser Pro Ser Arg Arg Thr Lys Ser Ser Ala Leu Lys Leu Ser Thr  
 1105 1110 1115 1120

Phe Met Pro Tyr Gln Asp Arg Gly Gly Gln Glu Pro Ala Gly Ala Gly  
 1125 1130 1135

Ser Pro Ser Pro Pro Glu Asp Arg Asn Thr Lys Thr Ala Pro Val Arg  
 1140 1145 1150

Leu Leu Pro Ser Tyr Ser Ala Phe Ser His Ser Ser His Asp Ser Cys  
 1155 1160 1165

Lys Asp Ser Ala Thr Leu Glu Glu Ile Pro Leu Thr Gln Thr Ser Asp  
 1170 1175 1180

Phe Pro Pro Ala Ala Thr Pro Ala Ser Ala Gln Thr Ala Lys Arg Glu  
 1185 1190 1195 1200

Ile Tyr Leu

(2) INFORMATION FOR SEQ ID NO:104:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 2789 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (ix) FEATURE:  
 (A) NAME/KEY: CDS  
 (B) LOCATION: 115..2622

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:104:

CGAAAGCCAT GTCGGACTCG TCGCCCAGCG CCCAAGCGCT AACCCGCTGA AAGTTTCTCA	60
CGGAAATCTC AGGGACGATC TGGACCCCGC TGAGAGGAAC TGCTTTTGAG TGAG ATG	117
	Met
	1
GTC CCA GAG GCC TGG AGG AGC GGA CTG GTA AGC ACC GGG AGG GTA GTG	165
Val Pro Glu Ala Trp Arg Ser Gly Leu Val Ser Thr Gly Arg Val Val	
	5 10 15
GGA GTT TTG CTT CTG CTT GGT GCC TTG AAC AAG GCT TCC ACG GTC ATT	213
Gly Val Leu Leu Leu Leu Gly Ala Leu Asn Lys Ala Ser Thr Val Ile	
	20 25 30
CAC TAT GAG ATC CCG GAG GAA AGA GAG AAG GGT TTC GCT GTG GGC AAC	261
His Tyr Glu Ile Pro Glu Glu Arg Glu Lys Gly Phe Ala Val Gly Asn	
	35 40 45

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GTG	GTC	GCG	AAC	CTT	GGT	TTG	GAT	CTC	GGT	AGC	CTC	TCA	GCC	CGC	AGG	309
Val	Val	Ala	Asn	Leu	Gly	Leu	Asp	Leu	Gly	Ser	Leu	Ser	Ala	Arg	Arg	
50					55					60					65	
TTC	CCG	GTG	GTG	TCT	GGA	GCT	AGC	CGA	AGA	TTC	TTT	GAG	GTG	AAC	CGG	357
Phe	Pro	Val	Val	Ser	Gly	Ala	Ser	Arg	Arg	Phe	Phe	Glu	Val	Asn	Arg	
				70					75					80		
GAG	ACC	GGA	GAG	ATG	TTT	GTG	AAC	GAC	CGT	CTG	GAT	CGA	GAG	GAG	CTG	405
Glu	Thr	Gly	Glu	Met	Phe	Val	Asn	Asp	Arg	Leu	Asp	Arg	Glu	Glu	Leu	
			85					90					95			
TGT	GGG	ACA	CTG	CCC	TCT	TGC	ACT	GTA	ACT	CTG	GAG	TTG	GTA	GTG	GAG	453
Cys	Gly	Thr	Leu	Pro	Ser	Cys	Thr	Val	Thr	Leu	Glu	Leu	Val	Val	Glu	
		100					105					110				
AAC	CCG	CTG	GAG	CTG	TTC	AGC	GTG	GAA	GTG	GTG	ATC	CAG	GAC	ATC	AAC	501
Asn	Pro	Leu	Glu	Leu	Phe	Ser	Val	Glu	Val	Val	Ile	Gln	Asp	Ile	Asn	
	115					120					125					
GAC	AAC	AAT	CCT	GCT	TTC	CCT	ACC	CAG	GAA	ATG	AAA	TTG	GAG	ATT	AGC	549
Asp	Asn	Asn	Pro	Ala	Phe	Pro	Thr	Gln	Glu	Met	Lys	Leu	Glu	Ile	Ser	
130					135					140					145	
GAG	GCC	GTG	GCT	CCG	GGG	ACG	CGC	TTT	CCG	CTC	GAG	AGC	GCG	CAC	GAT	597
Glu	Ala	Val	Ala	Pro	Gly	Thr	Arg	Phe	Pro	Leu	Glu	Ser	Ala	His	Asp	
				150					155					160		
CCC	GAT	CTG	GGA	AGC	AAC	TCT	TTA	CAA	ACC	TAT	GAG	CTG	AGC	CGA	AAT	645
Pro	Asp	Leu	Gly	Ser	Asn	Ser	Leu	Gln	Thr	Tyr	Glu	Leu	Ser	Arg	Asn	
			165					170					175			
GAA	TAC	TTT	GCG	CTT	CGC	GTG	CAG	ACG	CGG	GAG	GAC	AGC	ACC	AAG	TAC	693
Glu	Tyr	Phe	Ala	Leu	Arg	Val	Gln	Thr	Arg	Glu	Asp	Ser	Thr	Lys	Tyr	
		180					185					190				
GCG	GAG	CTG	GTG	TTG	GAG	CGC	GCC	CTG	GAC	CGA	GAA	CGG	GAG	CCT	AGT	741
Ala	Glu	Leu	Val	Leu	Glu	Arg	Ala	Leu	Asp	Arg	Glu	Arg	Glu	Pro	Ser	
	195					200					205					
CTC	CAG	TTA	GTG	CTG	ACG	GCG	TTG	GAC	GGA	GGG	ACC	CCA	GCT	CTC	TCC	789
Leu	Gln	Leu	Val	Leu	Thr	Ala	Leu	Asp	Gly	Gly	Thr	Pro	Ala	Leu	Ser	
210					215					220					225	
GCC	AGC	CTG	CCT	ATT	CAC	ATC	AAG	GTG	CTG	GAC	GCG	AAT	GAC	AAT	GCG	837
Ala	Ser	Leu	Pro	Ile	His	Ile	Lys	Val	Leu	Asp	Ala	Asn	Asp	Asn	Ala	
				230					235					240		
CCT	GTC	TTC	AAC	CAG	TCC	TTG	TAC	CGG	GCG	CGC	GTT	CCT	GGA	GGA	TGC	885
Pro	Val	Phe	Asn	Gln	Ser	Leu	Tyr	Arg	Ala	Arg	Val	Pro	Gly	Gly	Cys	
			245					250					255			
ACC	TCC	GGC	ACG	CGC	GTG	GTA	CAA	GTC	CTT	GCA	ACG	GAT	CTG	GAT	GAA	933
Thr	Ser	Gly	Thr	Arg	Val	Val	Gln	Val	Leu	Ala	Thr	Asp	Leu	Asp	Glu	
		260					265					270				
GGC	CCC	AAC	GGT	GAA	ATT	ATT	TAC	TCC	TTC	GGC	AGC	CAC	AAC	CGC	GCC	981
Gly	Pro	Asn	Gly	Glu	Ile	Ile	Tyr	Ser	Phe	Gly	Ser	His	Asn	Arg	Ala	
	275					280					285					

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GGC GTG CGG CAA CTA TTC GCC TTA GAC CTT GTA ACC GGG ATG CTG ACA Gly Val Arg Gln Leu Phe Ala Leu Asp Leu Val Thr Gly Met Leu Thr 290 295 300 305	1029
ATC AAG GGT CGG CTG GAC TTC GAG GAC ACC AAA CTC CAT GAG ATT TAC Ile Lys Gly Arg Leu Asp Phe Glu Asp Thr Lys Leu His Glu Ile Tyr 310 315 320	1077
ATC CAG GCC AAA GAC AAG GGC GCC AAT CCC GAA GGA GCA CAT TGC AAA Ile Gln Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys 325 330 335	1125
GTG TTG GTG GAG GTT GTG GAT GTG AAT GAC AAC GCC CCG GAG ATC ACA Val Leu Val Glu Val Val Asp Val Asn Asp Asn Ala Pro Glu Ile Thr 340 345 350	1173
GTC ACC TCC GTG TAC AGC CCA GTA CCC GAG GAT GCC TCT GGG ACT GTC Val Thr Ser Val Tyr Ser Pro Val Pro Glu Asp Ala Ser Gly Thr Val 355 360 365	1221
ATC GCT TTG CTC AGT GTG ACT GAC CTG GAT GCT GGC GAG AAC GGG CTG Ile Ala Leu Leu Ser Val Thr Asp Leu Asp Ala Gly Glu Asn Gly Leu 370 375 380 385	1269
GTG ACC TGC GAA GTT CCA CCG GGT CTC CCT TTC AGC CTT ACT TCT TCC Val Thr Cys Glu Val Pro Pro Gly Leu Pro Phe Ser Leu Thr Ser Ser 390 395 400	1317
CTC AAG AAT TAC TTC ACT TTG AAA ACC AGT GCA GAC CTG GAT CGG GAG Leu Lys Asn Tyr Phe Thr Leu Lys Thr Ser Ala Asp Leu Asp Arg Glu 405 410 415	1365
ACT GTG CCA GAA TAC AAC CTC AGC ATC ACC GCC CGA GAC GCC GGA ACC Thr Val Pro Glu Tyr Asn Leu Ser Ile Thr Ala Arg Asp Ala Gly Thr 420 425 430	1413
CCT TCC CTC TCA GCC CTT ACA ATA GTG CGT GTT CAA GTG TCC GAC ATC Pro Ser Leu Ser Ala Leu Thr Ile Val Arg Val Gln Val Ser Asp Ile 435 440 445	1461
AAT GAC AAC CCT CCA CAA TCT TCT CAA TCT TCC TAC GAC GTT TAC ATT Asn Asp Asn Pro Pro Gln Ser Ser Gln Ser Ser Tyr Asp Val Tyr Ile 450 455 460 465	1509
GAA GAA AAC AAC CTC CCC GGG GCT CCA ATA CTA AAC CTA AGT GTC TGG Glu Glu Asn Asn Leu Pro Gly Ala Pro Ile Leu Asn Leu Ser Val Trp 470 475 480	1557
GAC CCC GAC GCC CCG CAG AAT GCT CGG CTT TCT TTC TTT CTC TTG GAG Asp Pro Asp Ala Pro Gln Asn Ala Arg Leu Ser Phe Phe Leu Leu Glu 485 490 495	1605
CAA GGA GCT GAA ACC GGG CTA GTG GGT CGC TAT TTC ACA ATA AAT CGT Gln Gly Ala Glu Thr Gly Leu Val Gly Arg Tyr Phe Thr Ile Asn Arg 500 505 510	1653
GAC AAT GGC ATA GTG TCA TCC TTA GTG CCC CTA GAC TAT GAG GAT CGG Asp Asn Gly Ile Val Ser Ser Leu Val Pro Leu Asp Tyr Glu Asp Arg 515 520 525	1701

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CGG GAA TTT GAA TTA ACA GCT CAT ATC AGC GAT GGG GGC ACC CCG GTC Arg Glu Phe Glu Leu Thr Ala His Ile Ser Asp Gly Gly Thr Pro Val 530 535 540 545	1749
CTA GCC ACC AAC ATC AGC GTG AAC ATA TTT GTC ACT GAT CGC AAT GAC Leu Ala Thr Asn Ile Ser Val Asn Ile Phe Val Thr Asp Arg Asn Asp 550 555 560	1797
AAT GCC CCC CAG GTC CTA TAT CCT CGG CCA GGT GGG AGC TCG GTG GAG Asn Ala Pro Gln Val Leu Tyr Pro Arg Pro Gly Gly Ser Ser Val Glu 565 570 575	1845
ATG CTG CCT CGA GGT ACC TCA GCT GGC CAC CTA GTG TCA CGG GTG GTA Met Leu Pro Arg Gly Thr Ser Ala Gly His Leu Val Ser Arg Val Val 580 585 590	1893
GGC TGG GAC GCG GAT GCA GGG CAC AAT GCC TGG CTC TCC TAC AGT CTC Gly Trp Asp Ala Asp Ala Gly His Asn Ala Trp Leu Ser Tyr Ser Leu 595 600 605	1941
TTT GGA TCC CCT AAC CAG AGC CTT TTT GCC ATA GGG CTG CAC ACT GGT Phe Gly Ser Pro Asn Gln Ser Leu Phe Ala Ile Gly Leu His Thr Glu 610 615 620 625	1989
CAA ATC AGT ACT GCC CGT CCA GTC CAA GAC ACA GAT TCA CCC AGG CAG Gln Ile Ser Thr Ala Arg Pro Val Gln Asp Thr Asp Ser Pro Arg Gln 630 635 640	2037
ACT CTC ACT GTC TTG ATC AAA GAC AAT GGG GAG CCT TCG CTC TCC ACC Thr Leu Thr Val Leu Ile Lys Asp Asn Gly Glu Pro Ser Leu Ser Thr 645 650 655	2085
ACT GCT ACC CTC ACT GTG TCA GTA ACC GAG GAC TCT CCT GAA GCC CGA Thr Ala Thr Leu Thr Val Ser Val Thr Glu Asp Ser Pro Glu Ala Arg 660 665 670	2133
GCC GAG TTC CCC TCT GGC TCT GCC CCC CGG GAG CAG AAA AAA AAT CTC Ala Glu Phe Pro Ser Gly Ser Ala Pro Arg Glu Gln Lys Lys Asn Leu 675 680 685	2181
ACC TTT TAT CTA CTT CTT TCT CTA ATC CTG GTT TCT GTG GGC TTC GTG Thr Phe Tyr Leu Leu Leu Ser Leu Ile Leu Val Ser Val Gly Phe Val 690 695 700 705	2229
GTC ACA GTG TTC GGA GTA ATC ATA TTC AAA GTT TAC AAG TGG AAG CAG Val Thr Val Phe Gly Val Ile Ile Phe Lys Val Tyr Lys Trp Lys Gln 710 715 720	2277
TCT AGA GAC CTA TAC CGA GCC CCG GTG AGC TCA CTG TAC CGA ACA CCA Ser Arg Asp Leu Tyr Arg Ala Pro Val Ser Ser Leu Tyr Arg Thr Pro 725 730 735	2325
GGG CCC TCC TTG CAC GCG GAC GCC GTG CGG GGA GGC CTG ATG TCG CCG Gly Pro Ser Leu His Ala Asp Ala Val Arg Gly Gly Leu Met Ser Pro 740 745 750	2373
CAC CTT TAC CAT CAG GTG TAT CTC ACC ACG GAC TCC CGC CGC AGC GAC His Leu Tyr His Gln Val Tyr Leu Thr Thr Asp Ser Arg Arg Ser Asp 755 760 765	2421

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CCG CTG CTG AAG AAA CCT GGT GCA GCC AGT CCA CTG GCC AGC CGC CAG	2469
Pro Leu Leu Lys Lys Pro Gly Ala Ala Ser Pro Leu Ala Ser Arg Gln	
770 775 780 785	
AAC ACG CTG CGG AGC TGT GAT CCG GTG TTC TAT AGG CAG GTG TTG GGT	2517
Asn Thr Leu Arg Ser Cys Asp Pro Val Phe Tyr Arg Gln Val Leu Gly	
790 795 800	
GCA GAG AGC GCC CCT CCC GGA CAG GTA AGG TTT AGC AAG TCA TGC TTG	2565
Ala Glu Ser Ala Pro Pro Gly Gln Val Arg Phe Ser Lys Ser Cys Leu	
805 810 815	
ACC CTG TTA GTG CCT TTT TAT TCC TAC ATC ATA TTG AGA AGG CTG GAG	2613
Thr Leu Leu Val Pro Phe Tyr Ser Tyr Ile Ile Leu Arg Arg Leu Glu	
820 825 830	
CTG TTT TTT TAGTGATGAA GATGTTTTCC TGGTGATGCA TTCACACTTT	2662
Leu Phe Phe	
835	
CAACTGGCTC TTCCTAGATC AAAGTTAGTG CCTTTGTGAG ATGGTGGCCT GCCAGAGTGT	2722
GGTTTGTGGT CCCATTTCAG GGGGAAGATA CTTGACTCAT CTGTGGACCT AATTCACATC	2782
CTCAGCG	2789

(2) INFORMATION FOR SEQ ID NO:105:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 836 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:105:

Met Val Pro Glu Ala Trp Arg Ser Gly Leu Val Ser Thr Gly Arg Val	
1 5 10 15	
Val Gly Val Leu Leu Leu Leu Gly Ala Leu Asn Lys Ala Ser Thr Val	
20 25 30	
Ile His Tyr Glu Ile Pro Glu Glu Arg Glu Lys Gly Phe Ala Val Gly	
35 40 45	
Asn Val Val Ala Asn Leu Gly Leu Asp Leu Gly Ser Leu Ser Ala Arg	
50 55 60	
Arg Phe Pro Val Val Ser Gly Ala Ser Arg Arg Phe Phe Glu Val Asn	
65 70 75 80	
Arg Glu Thr Gly Glu Met Phe Val Asn Asp Arg Leu Asp Arg Glu Glu	
85 90 95	
Leu Cys Gly Thr Leu Pro Ser Cys Thr Val Thr Leu Glu Leu Val Val	
100 105 110	
Glu Asn Pro Leu Glu Leu Phe Ser Val Glu Val Val Ile Gln Asp Ile	
115 120 125	

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Asn Asp Asn Asn Pro Ala Phe Pro Thr Gln Glu Met Lys Leu Glu Ile  
 130 135 140  
 Ser Glu Ala Val Ala Pro Gly Thr Arg Phe Pro Leu Glu Ser Ala His  
 145 150 155 160  
 Asp Pro Asp Leu Gly Ser Asn Ser Leu Gln Thr Tyr Glu Leu Ser Arg  
 165 170 175  
 Asn Glu Tyr Phe Ala Leu Arg Val Gln Thr Arg Glu Asp Ser Thr Lys  
 180 185 190  
 Tyr Ala Glu Leu Val Leu Glu Arg Ala Leu Asp Arg Glu Arg Glu Pro  
 195 200 205  
 Ser Leu Gln Leu Val Leu Thr Ala Leu Asp Gly Gly Thr Pro Ala Leu  
 210 215 220  
 Ser Ala Ser Leu Pro Ile His Ile Lys Val Leu Asp Ala Asn Asp Asn  
 225 230 235 240  
 Ala Pro Val Phe Asn Gln Ser Leu Tyr Arg Ala Arg Val Pro Gly Gly  
 245 250 255  
 Cys Thr Ser Gly Thr Arg Val Val Gln Val Leu Ala Thr Asp Leu Asp  
 260 265 270  
 Glu Gly Pro Asn Gly Glu Ile Ile Tyr Ser Phe Gly Ser His Asn Arg  
 275 280 285  
 Ala Gly Val Arg Gln Leu Phe Ala Leu Asp Leu Val Thr Gly Met Leu  
 290 295 300  
 Thr Ile Lys Gly Arg Leu Asp Phe Glu Asp Thr Lys Leu His Glu Ile  
 305 310 315 320  
 Tyr Ile Gln Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys  
 325 330 335  
 Lys Val Leu Val Glu Val Val Asp Val Asn Asp Asn Ala Pro Glu Ile  
 340 345 350  
 Thr Val Thr Ser Val Tyr Ser Pro Val Pro Glu Asp Ala Ser Gly Thr  
 355 360 365  
 Val Ile Ala Leu Leu Ser Val Thr Asp Leu Asp Ala Gly Glu Asn Gly  
 370 375 380  
 Leu Val Thr Cys Glu Val Pro Pro Gly Leu Pro Phe Ser Leu Thr Ser  
 385 390 395 400  
 Ser Leu Lys Asn Tyr Phe Thr Leu Lys Thr Ser Ala Asp Leu Asp Arg  
 405 410 415  
 Glu Thr Val Pro Glu Tyr Asn Leu Ser Ile Thr Ala Arg Asp Ala Gly  
 420 425 430  
 Thr Pro Ser Leu Ser Ala Leu Thr Ile Val Arg Val Gln Val Ser Asp  
 435 440 445

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Ile Asn Asp Asn Pro Pro Gln Ser Ser Gln Ser Ser Tyr Asp Val Tyr  
450 455 460

Ile Glu Glu Asn Asn Leu Pro Gly Ala Pro Ile Leu Asn Leu Ser Val  
465 470 475 480

Trp Asp Pro Asp Ala Pro Gln Asn Ala Arg Leu Ser Phe Phe Leu Leu  
485 490 495

Glu Gln Gly Ala Glu Thr Gly Leu Val Gly Arg Tyr Phe Thr Ile Asn  
500 505 510

Arg Asp Asn Gly Ile Val Ser Ser Leu Val Pro Leu Asp Tyr Glu Asp  
515 520 525

Arg Arg Glu Phe Glu Leu Thr Ala His Ile Ser Asp Gly Gly Thr Pro  
530 535 540

Val Leu Ala Thr Asn Ile Ser Val Asn Ile Phe Val Thr Asp Arg Asn  
545 550 555 560

Asp Asn Ala Pro Gln Val Leu Tyr Pro Arg Pro Gly Gly Ser Ser Val  
565 570 575

Glu Met Leu Pro Arg Gly Thr Ser Ala Gly His Leu Val Ser Arg Val  
580 585 590

Val Gly Trp Asp Ala Asp Ala Gly His Asn Ala Trp Leu Ser Tyr Ser  
595 600 605

Leu Phe Gly Ser Pro Asn Gln Ser Leu Phe Ala Ile Gly Leu His Thr  
610 615 620

Gly Gln Ile Ser Thr Ala Arg Pro Val Gln Asp Thr Asp Ser Pro Arg  
625 630 635 640

Gln Thr Leu Thr Val Leu Ile Lys Asp Asn Gly Glu Pro Ser Leu Ser  
645 650 655

Thr Thr Ala Thr Leu Thr Val Ser Val Thr Glu Asp Ser Pro Glu Ala  
660 665 670

Arg Ala Glu Phe Pro Ser Gly Ser Ala Pro Arg Glu Gln Lys Lys Asn  
675 680 685

Leu Thr Phe Tyr Leu Leu Leu Ser Leu Ile Leu Val Ser Val Gly Phe  
690 695 700

Val Val Thr Val Phe Gly Val Ile Ile Phe Lys Val Tyr Lys Trp Lys  
705 710 715 720

Gln Ser Arg Asp Leu Tyr Arg Ala Pro Val Ser Ser Leu Tyr Arg Thr  
725 730 735

Pro Gly Pro Ser Leu His Ala Asp Ala Val Arg Gly Gly Leu Met Ser  
740 745 750

Pro His Leu Tyr His Gln Val Tyr Leu Thr Thr Asp Ser Arg Arg Ser  
755 760 765

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Asp Pro Leu Leu Lys Lys Pro Gly Ala Ala Ser Pro Leu Ala Ser Arg  
 770 775 780  
 Gln Asn Thr Leu Arg Ser Cys Asp Pro Val Phe Tyr Arg Gln Val Leu  
 785 790 795 800  
 Gly Ala Glu Ser Ala Pro Pro Gly Gln Val Arg Phe Ser Lys Ser Cys  
 805 810 815  
 Leu Thr Leu Leu Val Pro Phe Tyr Ser Tyr Ile Ile Leu Arg Arg Leu  
 820 825 830  
 Glu Leu Phe Phe  
 835

(2) INFORMATION FOR SEQ ID NO:106:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 2751 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

- (ix) FEATURE:  
 (A) NAME/KEY: CDS  
 (B) LOCATION: 115..2160

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:106:

CGAAAGCCAT GTCGGACTCG TCGCCCAGCG CCCAAGCGCT AACCCGCTGA AAGTTTCTCA	60
GCGAAATCTC AGGGACGATC TGGACCCCGC TGAGAGGAAC TGCTTTTGAG TGAG ATG	117
	Met
	1
GTC CCA GAG GCC TGG AGG AGC GGA CTG GTA AGC ACC GGG AGG GTA GTG	165
Val Pro Glu Ala Trp Arg Ser Gly Leu Val Ser Thr Gly Arg Val Val	
5 10 15	
GGA GTT TTG CTT CTG CTT GGT GCC TTG AAC AAG GCT TCC ACG GTC ATT	213
Gly Val Leu Leu Leu Leu Gly Ala Leu Asn Lys Ala Ser Thr Val Ile	
20 25 30	
CAC TAT GAG ATC CCG GAG GAA AGA GAG AAG GGT TTC GCT GTG GGC AAC	261
His Tyr Glu Ile Pro Glu Glu Arg Glu Lys Gly Phe Ala Val Gly Asn	
35 40 45	
GTG GTC GCG AAC CTT GGT TTG GAT CTC GGT AGC CTC TCA GCC CGC AGG	309
Val Val Ala Asn Leu Gly Leu Asp Leu Gly Ser Leu Ser Ala Arg Arg	
50 55 60 65	
TTC CCG GTG GTG TCT GGA GCT AGC CGA AGA TTC TTT GAG GTG AAC CGG	357
Phe Pro Val Val Ser Gly Ala Ser Arg Arg Phe Phe Glu Val Asn Arg	
70 75 80	

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GAG ACC GGA GAG ATG TTT GTG AAC GAC CGT CTG GAT CGA GAG GAG CTG	405
Glu Thr Gly Glu Met Phe Val Asn Asp Arg Leu Asp Arg Glu Glu Leu	
85 90 95	
TGT GGG ACA CTG CCC TCT TGC ACT GTA ACT CTG GAG TTG GTA GTG GAG	453
Cys Gly Thr Leu Pro Ser Cys Thr Val Thr Leu Glu Leu Val Val Glu	
100 105 110	
AAC CCG CTG GAG CTG TTC AGC GTG GAA GTG GTG ATC CAG GAC ATC AAC	501
Asn Pro Leu Glu Leu Phe Ser Val Glu Val Val Ile Gln Asp Ile Asn	
115 120 125	
GAC AAC AAT CCT GCT TTC CCT ACC CAG GAA ATG AAA TTG GAG ATT AGC	549
Asp Asn Asn Pro Ala Phe Pro Thr Gln Glu Met Lys Leu Glu Ile Ser	
130 135 140 145	
GAG GCC GTG GCT CCG GGG ACG CGC TTT CCG CTC GAG AGC GCG CAC GAT	597
Glu Ala Val Ala Pro Gly Thr Arg Phe Pro Leu Glu Ser Ala His Asp	
150 155 160	
CCC GAT CTG GGA AGC AAC TCT TTA CAA ACC TAT GAG CTG AGC CGA AAT	645
Pro Asp Leu Gly Ser Asn Ser Leu Gln Thr Tyr Glu Leu Ser Arg Asn	
165 170 175	
GAA TAC TTT GCG CTT CGC GTG CAG ACG CGG GAG GAC AGC ACC AAG TAC	693
Glu Tyr Phe Ala Leu Arg Val Gln Thr Arg Glu Asp Ser Thr Lys Tyr	
180 185 190	
GCG GAG CTG GTG TTG GAG CGC GCC CTG GAC CGA GAA CGG GAG CCT AGT	741
Ala Glu Leu Val Leu Glu Arg Ala Leu Asp Arg Glu Arg Glu Pro Ser	
195 200 205	
CTC CAG TTA GTG CTG ACG GCG TTG GAC GGA GGG ACC CCA GCT CTC TCC	789
Leu Gln Leu Val Leu Thr Ala Leu Asp Gly Gly Thr Pro Ala Leu Ser	
210 215 220 225	
GCC AGC CTG CCT ATT CAC ATC AAG GTG CTG GAC GCG AAT GAC AAT GCG	837
Ala Ser Leu Pro Ile His Ile Lys Val Leu Asp Ala Asn Asp Asn Ala	
230 235 240	
CCT GTC TTC AAC CAG TCC TTG TAC CGG GCG CGC GTT CCT GGA GGA TGC	885
Pro Val Phe Asn Gln Ser Leu Tyr Arg Ala Arg Val Pro Gly Gly Cys	
245 250 255	
ACC TCC GGC ACG CGC GTG GTA CAA GTC CTT GCA ACG GAT CTG GAT GAA	933
Thr Ser Gly Thr Arg Val Val Gln Val Leu Ala Thr Asp Leu Asp Glu	
260 265 270	
GGC CCC AAC GGT GAA ATT ATT TAC TCC TTC GGC AGC CAC AAC CGC GCC	981
Gly Pro Asn Gly Glu Ile Ile Tyr Ser Phe Gly Ser His Asn Arg Ala	
275 280 285	
GGC GTG CGG CAA CTA TTC GCC TTA GAC CTT GTA ACC GGG ATG CTG ACA	1029
Gly Val Arg Gln Leu Phe Ala Leu Asp Leu Val Thr Gly Met Leu Thr	
290 295 300 305	
ATC AAG GGT CGG CTG GAC TTC GAG GAC ACC AAA CTC CAT GAG ATT TAC	1077
Ile Lys Gly Arg Leu Asp Phe Glu Asp Thr Lys Leu His Glu Ile Tyr	
310 315 320	

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ATC CAG GCC AAA GAC AAG GGC GCC AAT CCC GAA GGA GCA CAT TGC AAA Ile Gln Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys 325 330 335	1125
GTG TTG GTG GAG GTT GTG GAT GTG AAT GAC AAC GCC CCG GAG ATC ACA Val Leu Val Glu Val Val Asp Val Asn Asp Asn Ala Pro Glu Ile Thr 340 345 350	1173
GTC ACC TCC GTG TAC AGC CCA GTA CCC GAG GAT GCC TCT GGG ACT GTC Val Thr Ser Val Tyr Ser Pro Val Pro Glu Asp Ala Ser Gly Thr Val 355 360 365	1221
ATC GCT TTG CTC AGT GTG ACT GAC CTG GAT GCT GGC GAG AAC GGG CTG Ile Ala Leu Leu Ser Val Thr Asp Leu Asp Ala Gly Glu Asn Gly Leu 370 375 380 385	1269
GTG ACC TGC GAA GTT CCA CCG GGT CTC CCT TTC AGC CTT ACT TCT TCC Val Thr Cys Glu Val Pro Pro Gly Leu Pro Phe Ser Leu Thr Ser Ser 390 395 400	1317
CTC AAG AAT TAC TTC ACT TTG AAA ACC AGT GCA GAC CTG GAT CGG GAG Leu Lys Asn Tyr Phe Thr Leu Lys Thr Ser Ala Asp Leu Asp Arg Glu 405 410 415	1365
ACT GTG CCA GAA TAC AAC CTC AGC ATC ACC GCC CGA GAC GCC GGA ACC Thr Val Pro Glu Tyr Asn Leu Ser Ile Thr Ala Arg Asp Ala Gly Thr 420 425 430	1413
CCT TCC CTC TCA GCC CTT ACA ATA GTG CGT GTT CAA GTG TCC GAC ATC Pro Ser Leu Ser Ala Leu Thr Ile Val Arg Val Gln Val Ser Asp Ile 435 440 445	1461
AAT GAC AAC CCT CCA CAA TCT TCT CAA TCT TCC TAC GAC GTT TAC ATT Asn Asp Asn Pro Pro Gln Ser Ser Gln Ser Ser Tyr Asp Val Tyr Ile 450 455 460 465	1509
GAA GAA AAC AAC CTC CCC GGG GCT CCA ATA CTA AAC CTA AGT GTC TGG Glu Glu Asn Asn Leu Pro Gly Ala Pro Ile Leu Asn Leu Ser Val Trp 470 475 480	1557
GAC CCC GAC GCC CCG CAG AAT GCT CGG CTT TCT TTC TTT CTC TTG GAG Asp Pro Asp Ala Pro Gln Asn Ala Arg Leu Ser Phe Phe Leu Leu Glu 485 490 495	1605
CAA GGA GCT GAA ACC GGG CTA GTG GGT CGC TAT TTC ACA ATA AAT CGT Gln Gly Ala Glu Thr Gly Leu Val Gly Arg Tyr Phe Thr Ile Asn Arg 500 505 510	1653
GAC AAT GGC ATA GTG TCA TCC TTA GTG CCC CTA GAC TAT GAG GAT CGG Asp Asn Gly Ile Val Ser Ser Leu Val Pro Leu Asp Tyr Glu Asp Arg 515 520 525	1701
CGG GAA TTT GAA TTA ACA GCT CAT ATC AGC GAT GGG GGC ACC CCG GTC Arg Glu Phe Glu Leu Thr Ala His Ile Ser Asp Gly Gly Thr Pro Val 530 535 540 545	1749
CTA GCC ACC AAC ATC AGC GTG AAC ATA TTT GTC ACT GAT CGC AAT GAC Leu Ala Thr Asn Ile Ser Val Asn Ile Phe Val Thr Asp Arg Asn Asp 550 555 560	1797

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AAT GCC CCC CAG GTC CTA TAT CCT CGG CCA GGT GGG AGC TCG GTG GAG Asn Ala Pro Gln Val Leu Tyr Pro Arg Pro Gly Gly Ser Ser Val Glu 565 570 575	1845
ATG CTG CCT CGA GGT ACC TCA GCT GGC CAC CTA GTG TCA CGG GTG GTA Met Leu Pro Arg Gly Thr Ser Ala Gly His Leu Val Ser Arg Val Val 580 585 590	1893
GGC TGG GAC GCG GAT GCA GGG CAC AAT GCC TGG CTC TCC TAC AGT CTC Gly Trp Asp Ala Asp Ala Gly His Asn Ala Trp Leu Ser Tyr Ser Leu 595 600 605	1941
TTT GGA TCC CCT AAC CAG AGC CTT TTT GCC ATA GGG CTG CAC ACT GGT Phe Gly Ser Pro Asn Gln Ser Leu Phe Ala Ile Gly Leu His Thr Gly 610 615 620 625	1989
CAA ATC AGT ACT GCC CGT CCA GTC CAA GAC ACA GAT TCA CCC AGG CAG Gln Ile Ser Thr Ala Arg Pro Val Gln Asp Thr Asp Ser Pro Arg Gln 630 635 640	2037
ACT CTC ACT GTC TTG ATC AAA GAC AAT GGG GAG CCT TCG CTC TCC ACC Thr Leu Thr Val Leu Ile Lys Asp Asn Gly Glu Pro Ser Leu Ser Thr 645 650 655	2085
ACT GCT ACC CTC ACT GTG TCA GTA ACC GAG GAC TCT CCT GAA GCC CGA Thr Ala Thr Leu Thr Val Ser Val Thr Glu Asp Ser Pro Glu Ala Arg 660 665 670	2133
GCC GAG TTC CCC TCT GGC TCT GCC AGT TAAACCTTCT TTAATTATGG Ala Glu Phe Pro Ser Gly Ser Ala Ser 675 680	2180
ATTAGCCATT AACATTTTTG AAACGTGGAC CATTTAACCT CGGCCTACCC CCTCCAAC TG	2240
TCCTGGTGAT GAGTTCATTA GCTAAGTTAA ATTAATTGAA CTTTGATCTA AACCAAAACA	2300
AATCAGGAAA ATAAAGCTGT AAAGGAAC TT ATCAAGCATT CCAAAACCAA CTAGAAATTA	2360
CTTGAAGTTT CGAGTGAGCA TTGCCTGTGC CAGTATTCTT CATTATAGGA TTATAAACTC	2420
GTTTTTTTTCC CAAAGCGCAT GTCTACGCCA GGCAGAGGAG TAATTATTCA GCCAATTTCA	2480
TGGATGTAAC GATGGATATA AATAATTGAT AGCACCTAGA GGCTTCCAGT TTGGGTGGAA	2540
GGCTAAAAGT AGAGGGGAAC TCACTCACTT GAGAAATGAT ATTTAAGTGA ATAAATAGTT	2600
CTCTTCTATG AAAC TATTAC TATTTAGTTC TCTGGAAAAC TTAAGTGTAT TAATGATTAG	2660
AACATCAAAT CCTAAGTAAA GAAATGACAT TTAAATATA AAAAGCCAAA CTTTAAATAA	2720
ATCATAGAGA CCTCAGACAT AATATAGGAA A	2751

(2) INFORMATION FOR SEQ ID NO:107:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 682 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:107:

Met	Val	Pro	Glu	Ala	Trp	Arg	Ser	Gly	Leu	Val	Ser	Thr	Gly	Arg	Val	1	5	10	15
Val	Gly	Val	Leu	Leu	Leu	Leu	Gly	Ala	Leu	Asn	Lys	Ala	Ser	Thr	Val	20	25	30	
Ile	His	Tyr	Glu	Ile	Pro	Glu	Glu	Arg	Glu	Lys	Gly	Phe	Ala	Val	Gly	35	40	45	
Asn	Val	Val	Ala	Asn	Leu	Gly	Leu	Asp	Leu	Gly	Ser	Leu	Ser	Ala	Arg	50	55	60	
Arg	Phe	Pro	Val	Val	Ser	Gly	Ala	Ser	Arg	Arg	Phe	Phe	Glu	Val	Asn	65	70	75	80
Arg	Glu	Thr	Gly	Glu	Met	Phe	Val	Asn	Asp	Arg	Leu	Asp	Arg	Glu	Glu	85	90	95	
Leu	Cys	Gly	Thr	Leu	Pro	Ser	Cys	Thr	Val	Thr	Leu	Glu	Leu	Val	Val	100	105	110	
Glu	Asn	Pro	Leu	Glu	Leu	Phe	Ser	Val	Glu	Val	Val	Ile	Gln	Asp	Ile	115	120	125	
Asn	Asp	Asn	Asn	Pro	Ala	Phe	Pro	Thr	Gln	Glu	Met	Lys	Leu	Glu	Ile	130	135	140	
Ser	Glu	Ala	Val	Ala	Pro	Gly	Thr	Arg	Phe	Pro	Leu	Glu	Ser	Ala	His	145	150	155	160
Asp	Pro	Asp	Leu	Gly	Ser	Asn	Ser	Leu	Gln	Thr	Tyr	Glu	Leu	Ser	Arg	165	170	175	
Asn	Glu	Tyr	Phe	Ala	Leu	Arg	Val	Gln	Thr	Arg	Glu	Asp	Ser	Thr	Lys	180	185	190	
Tyr	Ala	Glu	Leu	Val	Leu	Glu	Arg	Ala	Leu	Asp	Arg	Glu	Arg	Glu	Pro	195	200	205	
Ser	Leu	Gln	Leu	Val	Leu	Thr	Ala	Leu	Asp	Gly	Gly	Thr	Pro	Ala	Leu	210	215	220	
Ser	Ala	Ser	Leu	Pro	Ile	His	Ile	Lys	Val	Leu	Asp	Ala	Asn	Asp	Asn	225	230	235	240
Ala	Pro	Val	Phe	Asn	Gln	Ser	Leu	Tyr	Arg	Ala	Arg	Val	Pro	Gly	Gly	245	250	255	
Cys	Thr	Ser	Gly	Thr	Arg	Val	Val	Gln	Val	Leu	Ala	Thr	Asp	Leu	Asp	260	265	270	
Glu	Gly	Pro	Asn	Gly	Glu	Ile	Ile	Tyr	Ser	Phe	Gly	Ser	His	Asn	Arg	275	280	285	
Ala	Gly	Val	Arg	Gln	Leu	Phe	Ala	Leu	Asp	Leu	Val	Thr	Gly	Met	Leu	290	295	300	
Thr	Ile	Lys	Gly	Arg	Leu	Asp	Phe	Glu	Asp	Thr	Lys	Leu	His	Glu	Ile	305	310	315	320

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Tyr Ile Gln Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys  
 325 330 335  
 Lys Val Leu Val Glu Val Val Asp Val Asn Asp Asn Ala Pro Glu Ile  
 340 345 350  
 Thr Val Thr Ser Val Tyr Ser Pro Val Pro Glu Asp Ala Ser Gly Thr  
 355 360 365  
 Val Ile Ala Leu Leu Ser Val Thr Asp Leu Asp Ala Gly Glu Asn Gly  
 370 375 380  
 Leu Val Thr Cys Glu Val Pro Pro Gly Leu Pro Phe Ser Leu Thr Ser  
 385 390 395 400  
 Ser Leu Lys Asn Tyr Phe Thr Leu Lys Thr Ser Ala Asp Leu Asp Arg  
 405 410 415  
 Glu Thr Val Pro Glu Tyr Asn Leu Ser Ile Thr Ala Arg Asp Ala Gly  
 420 425 430  
 Thr Pro Ser Leu Ser Ala Leu Thr Ile Val Arg Val Gln Val Ser Asp  
 435 440 445  
 Ile Asn Asp Asn Pro Pro Gln Ser Ser Gln Ser Ser Tyr Asp Val Tyr  
 450 455 460  
 Ile Glu Glu Asn Asn Leu Pro Gly Ala Pro Ile Leu Asn Leu Ser Val  
 465 470 475 480  
 Trp Asp Pro Asp Ala Pro Gln Asn Ala Arg Leu Ser Phe Phe Leu Leu  
 485 490 495  
 Glu Gln Gly Ala Glu Thr Gly Leu Val Gly Arg Tyr Phe Thr Ile Asn  
 500 505 510  
 Arg Asp Asn Gly Ile Val Ser Ser Leu Val Pro Leu Asp Tyr Glu Asp  
 515 520 525  
 Arg Arg Glu Phe Glu Leu Thr Ala His Ile Ser Asp Gly Gly Thr Pro  
 530 535 540  
 Val Leu Ala Thr Asn Ile Ser Val Asn Ile Phe Val Thr Asp Arg Asn  
 545 550 555 560  
 Asp Asn Ala Pro Gln Val Leu Tyr Pro Arg Pro Gly Gly Ser Ser Val  
 565 570 575  
 Glu Met Leu Pro Arg Gly Thr Ser Ala Gly His Leu Val Ser Arg Val  
 580 585 590  
 Val Gly Trp Asp Ala Asp Ala Gly His Asn Ala Trp Leu Ser Tyr Ser  
 595 600 605  
 Leu Phe Gly Ser Pro Asn Gln Ser Leu Phe Ala Ile Gly Leu His Thr  
 610 615 620  
 Gly Gln Ile Ser Thr Ala Arg Pro Val Gln Asp Thr Asp Ser Pro Arg  
 625 630 635 640

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Gln Thr Leu Thr Val Leu Ile Lys Asp Asn Gly Glu Pro Ser Leu Ser  
645 650 655  
Thr Thr Ala Thr Leu Thr Val Ser Val Thr Glu Asp Ser Pro Glu Ala  
660 665 670  
Arg Ala Glu Phe Pro Ser Gly Ser Ala Ser  
675 680

(2) INFORMATION FOR SEQ ID NO:108:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2831 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:108:

GAATTCGGCA CGAGGCTGAA CTGAGGGTGA CGGACATAAA CGACTATTCT CCAGTGTTCA	60
GTGAAAGAGA AATGATACTG AGGATACCAG AAAACAGTGC TCGGGGAAAT ACATTCCCTT	120
TAAACAATGC TCTGGACTCA GACGTAGATA TCAACAATAT CCAGACCTAT AGGCTCAGCT	180
CAAACCTCTCA TTTCCTGGTT GTAACCCGCA ACCGCAGTGA TGGCAGGAAG TACCCAGAGC	240
TGGTGCTGGA GAAAGAACTG GATCGAGAGG AGGAACCTGA GCTGAGGTTA ACGCTGACAG	300
CTTTGGATGG TGGCTCTCCT CCCCAGTCTG GGACGACACA GGTCTTCATT GAAGTAGTGG	360
ACACCAACGA TAATGCACCC GAGTTTCAGC AGCCAACATA CCAAGTGCAA ACTCCCGAGA	420
ACAGTCCCAC CGGCTCTCTG GTACTCACAG TCTCAGCCAA TGACTTAGAC AGTGGAGACT	480
ATGGGAAAGT CTTGTACGCA CTTTCGCAAC CCTCAGAAGA TATTAGCAAA ACATTTCGAGG	540
TAAACCCTGT AACCGGGGAA ATTCGCCTAC GAAAAGAGGT GAATTTTGAA ACTATTCCTT	600
CGTATGAAGT GGTATCAAG GGGACGGACG GGGGAGGTCT CTCAGGAAAA TGCCTCTGT	660
TACTGCAGGT GGTGGACGTG AATGACAATG CCCCAGAAGT GATGCTATCT GCGCTAACCA	720
ACCCAGTCCC AGAAAATTCC CCCGATGAGG TAGTGGCTGT TTTCAGTGTT AGAGATCCTG	780
ACTCTGGGAA CAACGGAAAA GTGATTGCAT CCATCGAGGA AGACCTGCCC TTTCTTCTAA	840
AATCTTCAGG AAAGAACTTT TACACTTTAG TAACCAAGGG AGCACTTGAC AGGGAAGAAA	900
GAGAGCAATT GAACATCACC ATCACAGTCA CTGACCTGGG CATACCCAGG CTCACCACCC	960
AACACACCAT AACAGTGCAG GTGGCAGACA TCAACGACAA TGCCCCCTCC TTCACCCAAA	1020
CCTCCTACAC CATGTTTGTC CGCGAGAACA ACAGCCCCGC CCTGCACATA GGCACCATCA	1080
GCGCCACAGA CTCAGACTCA GGATCCAATG CCCACATCAC CTACTCGCTG CTACCGCCCC	1140

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AAGACCCACA	GCTGGCCCTC	GACTCGCTCA	TCTCCATCAA	TGTAGACAAC	GGGCAGCTGT	1200
TCGCGCTCAG	GGCGCTAGAC	TATGAGGCTC	TGCAGGGCTT	CGAGTTCCAT	GTGGGCGCCA	1260
CAGACCAAGG	CTCGCCCGCG	CTCAGCAGCC	AGGCTCTGGT	GCACGTGGTG	GTGTTGGACG	1320
ACAATGACAA	TGCGCCCTTC	GTGCTCTACC	CGCTGCAAAA	CGCCTCTGCA	CCCTTCACTG	1380
AGCTGCTGCC	CAGGGCGGCA	GAGCCTGGAT	ACCTGGTTAC	CAAGGTGGTA	GCTGTGGACC	1440
GCGACTCTGG	CCAGAATGCC	TGGCTGTCA	TCCAGCTGCT	CAAGGCCACG	GAGCCCGGGC	1500
TGTTCAACGT	ATGGGCGCAC	AATGGCGAGG	TACGCACCTC	CAGGCTGCTG	AGCGAGCGCG	1560
ACGCACCCAA	GCACAAGCTG	CTGCTGTTGG	TCAAGGACAA	TGGAGATCCT	CCACGCTCTG	1620
CCAGTGTTAC	TCTGCACGTG	CTAGTGGTGG	ATGCCTTCTC	TCAGCCCTAC	CTGCCTCTGC	1680
CAGAGGTGGC	GCACGACCCT	GCACAAGAAG	AAGATGCGCT	AACACTCTAC	CTGGTCATAG	1740
CTTTGGCATC	TGTGTCTTCT	CTCTTCTCT	TGTCTGTGCT	GCTGTTCGTG	GGGGTGAGGC	1800
TCTGCAGGAG	GGCCAGGGCA	GCCTCTCTGA	GTGCCTATTC	TGTGCCTGAA	GGCCACTTTC	1860
CTGGCCAGCT	GGTGGATGTC	AGAGGTATGG	GGACCCTGTC	CCAGAGCTAC	CAGTATGATG	1920
TATGTCTGAT	GGGGGATTCT	TCTGGGACCA	GCGAATTTAA	CTTCTTAAAG	CCAGTTCTGC	1980
CTAGCTCTCT	GCACCAGTGC	TCTGGGAAAG	AAATAGAGGA	AAATTCCACA	CTCCAGAATA	2040
GTTTTGGGTT	TCATCATTAA	TAGAAAATA	CTTTACAGAT	ATTTAATTCC	AAATATCATC	2100
TTGTTGATTA	ACTAAAGTCT	GTTACATGT	AGCTAGCTAG	CAACGATTTT	AATGTTCACT	2160
TTACCCATCT	TTTTTCAGGG	TCATGTCTAA	AGCTACAAGT	TTGNCTTTAC	TTATACTTGT	2220
CGCACAGAAT	NNNNNNNNNN	TGGTGTATAA	GTCACAGTCA	TGGGATACTG	GCACAAGATG	2280
GCAGCTTGAT	TGCTCAGTTA	TGGCTGCAAA	GGGGNGCTTG	AGTTTAGGGA	ATGTGTTAGA	2340
GCTGGAATAA	GTTTTCTGAG	AAATGTGTAA	GACAAATTTT	TTTTGCACAT	TCCCTGTGTT	2400
CCTGTACCCC	TGTTTCCAGA	ACTACGAAAT	GTGTCATCAG	AAGGCATGCT	CACATTTTCC	2460
CCTTTGTTTG	CGTGACCCGG	GTGCCAGAAA	TTAAATAAAA	TTAGCATGGA	GTTCAATGCA	2520
GCATTAAAAAC	AAAGTTACTT	CTACAAACCT	TTTATTTCGAC	GGTTAAAATT	GTAACCTCCC	2580
CACCCATGAG	GCTGGCTGTA	AGAACCAGTA	TGAATGGGTG	TCTATCGCAA	CCTTATTTTC	2640
AAAAATCAAA	CAAAAGGAGA	AATGAGAGAC	CAAACAACAC	GCTACAGGAA	AGATTTTCATA	2700
AGGATGTATG	TATGGACACA	AAACTGGGA	TACAGACATT	TTAAATCTGT	TGGTACCACA	2760
TGGTGGCGCT	GCAGGCTAAA	GAAATGCAAG	GGAAATTAAA	AAGAGGCTGA	GCTAGAAGTC	2820
AAAAAAAAAA	A					2831

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(2) INFORMATION FOR SEQ ID NO:109:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 3353 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 763..3123

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:109:

GTATTTTCC ACAGTTTAAA ATTTTCATAA AATCATAACT CTCTGACTTT ATGTAGAAAG	60
GATACCACAC TGAATTAAC GTGTAGCTTT TTCTTGATGT AATCCAACCA ATGGGAGCAC	120
AATTCTGGTA CATAGGCTGT CTAGAATTTG AAAGAAATTA AAGAATTCAT TTTGTTTTGC	180
TGATAAATTT TTAAGAAATC ACGTGGCTTT ATGTTATTAT TATTACAAGA TGA CTGATCA	240
CTATTATGTC TTCTTTCACT TCTCAATTTT CCTCAGAACA CTACACCCAG ACTACAGGCT	300
CTGGAGGGTG GGGACCATGT CTGGGTTGTT TACTGATGTA TTTCATAATT TGGCACATAG	360
AGACCAATAA TACTCCTTTA AATGAAGAAA TTAATAATTA CCATTGCGTG ATATTGTGAT	420
TACATCATTT CCTCCCAATT TCCAACTCC TAATAGAATA GAGAATAGAT CAATTGTAGC	480
AATTCGTTTC GAAGCAAAGA CAACGCATGG TGGCGCTGCA GGCTAAGGCT TCAAAAAAAG	540
GAAAAGGAAA AAGCCCATGA AATGCTACTA GCTACTTCAG ACCTCTTTCA GCCTAAGAGG	600
AAAGCCTGTT AGCAGAGCAC GGACCAAGTGT CTCCGGAGAA TGCTATTCTC CTACATTTC	660
GAACAGGTTA TCAACGCACA GATCGATCAC TGCCTCTGTC CCATCGCTCC CTGAAGTAGC	720
TCTGACTCCG GTTCCTTGAA AGGGGCGTGT ACAGAAGTAA AG ATG GAG CCT GCA	774
Met Glu Pro Ala	
1	
GGG GAG CGC TTT CCC GAA CAA AGG CAA GTC CTG ATT CTC CTT CTT TTA	822
Gly Glu Arg Phe Pro Glu Gln Arg Gln Val Leu Ile Leu Leu Leu Leu	
5 10 15 20	
CTG GAA GTG ACT CTG GCA GGC TGG GAA CCC CGT CGC TAT TCT GTG ATG	870
Leu Glu Val Thr Leu Ala Gly Trp Glu Pro Arg Arg Tyr Ser Val Met	
25 30 35	
GAG GAA ACA GAG AGA GGT TCT TTT GTA GCC AAC CTG GCC AAT GAC CTA	918
Glu Glu Thr Glu Arg Gly Ser Phe Val Ala Asn Leu Ala Asn Asp Leu	
40 45 50	
GGG CTG GGA GTG GGG GAG CTA GCC GAG CGG GGA GCC CGG GTA GTT TCT	966
Gly Leu Gly Val Gly Glu Leu Ala Glu Arg Gly Ala Arg Val Val Ser	
55 60 65	

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GAG GAT AAC GAA CAA GGC TTG CAG CTT GAT CTG CAG ACC GGG CAG TTG Glu Asp Asn Glu Gln Gly Leu Gln Leu Asp Leu Gln Thr Gly Gln Leu 70 75 80	1014
ATA TTA AAT GAG AAG CTG GAC CGG GAG AAG CTG TGT GGC CCT ACT GAG Ile Leu Asn Glu Lys Leu Asp Arg Glu Lys Leu Cys Gly Pro Thr Glu 85 90 95 100	1062
CCC TGT ATA ATG CAT TTC CAA GTG TTA CTG AAA AAA CCT TTG GAA GTA Pro Cys Ile Met His Phe Gln Val Leu Lys Lys Pro Leu Glu Val 105 110 115	1110
TTT CGA GCT GAA CTA CTA GTG ACA GAC ATA AAC GAT CAT TCT CCT GAG Phe Arg Ala Leu Leu Val Thr Asp Ile Asn Asp His Ser Pro Glu 120 125 130	1158
TTT CCT GAA AGA GAA ATG ACC CTG AAA ATC CCA GAA ACT AGC TCC CTT Phe Pro Glu Arg Glu Met Thr Leu Lys Ile Pro Glu Thr Ser Ser Leu 135 140 145	1206
GGG ACT GTG TTT CCT CTG AAA AAA GCT CGG GAC TTG GAC GTG GGC AGC Gly Thr Val Phe Pro Leu Lys Lys Ala Arg Asp Leu Asp Val Gly Ser 150 155 160	1254
AAT AAT GTT CAA AAC TAC AAT ATT TCT CCC AAT TCT CAT TTC CAT GTT Asn Asn Val Gln Asn Tyr Asn Ile Ser Pro Asn Ser His Phe His Val 165 170 175 180	1302
TCC ACT CGC ACC CGA GGG GAT GGC AGG AAA TAC CCA GAG CTG GTG CTG Ser Thr Arg Thr Arg Gly Asp Gly Arg Lys Tyr Pro Glu Leu Val Leu 185 190 195	1350
GAC ACA GAA CTG GAT CGC GAG GAG CAG GCC GAG CTC AGA TTA ACC TTG Asp Thr Glu Leu Asp Arg Glu Glu Gln Ala Glu Leu Arg Leu Thr Leu 200 205 210	1398
ACA GCG GTG GAC GGT GGC TCT CCA CCC CGA TCT GGC ACC GTC CAG ATC Thr Ala Val Asp Gly Gly Ser Pro Pro Arg Ser Gly Thr Val Gln Ile 215 220 225	1446
CTC ATC TTG GTC TTG GAC GCC AAT GAC AAT GCC CCG GAG TTT GTG CAG Leu Ile Leu Val Leu Asp Ala Asn Asp Asn Ala Pro Glu Phe Val Gln 230 235 240	1494
GCG CTC TAC GAG GTG CAG GTC CCA GAG AAC AGC CCA GTA GGC TCC CTA Ala Leu Tyr Glu Val Gln Val Pro Glu Asn Ser Pro Val Gly Ser Leu 245 250 255 260	1542
GTT GTC AAG GTC TCT GCT AGG GAT TTA GAC ACT GGG ACA AAT GGA GAG Val Val Lys Val Ser Ala Arg Asp Leu Asp Thr Gly Thr Asn Gly Glu 265 270 275	1590
ATA TCA TAC TCC CTT TAT TAC AGC TCT CAG GAG ATA GAC AAA CCT TTT Ile Ser Tyr Ser Leu Tyr Tyr Ser Ser Gln Glu Ile Asp Lys Pro Phe 280 285 290	1638
GAG CTA AGC AGC CTT TCA GGA GAA ATT CGA CTA ATT AAA AAA CTA GAT Glu Leu Ser Ser Leu Ser Gly Glu Ile Arg Leu Ile Lys Lys Leu Asp 295 300 305	1686

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TTT GAG ACA ATG TCT TCA TAT GAT CTA GAT ATA GAG GCA TCT GAT GGC Phe Glu Thr Met Ser Ser Tyr Asp Leu Asp Ile Glu Ala Ser Asp Gly 310 315 320	1734
GGG GGA CTT TCT GGA AAA TGC TCT GTC TCT GTT AAG GTG CTG GAT GTT Gly Gly Leu Ser Gly Lys Cys Ser Val Ser Val Lys Val Leu Asp Val 325 330 335 340	1782
AAC GAT AAC TTC CCG GAA CTA AGT ATT TCA TCA CTT ACC AGC CCT ATT Asn Asp Asn Phe Pro Glu Leu Ser Ile Ser Ser Leu Thr Ser Pro Ile 345 350 355	1830
CCC GAG AAT TCT CCA GAG ACA GAA GTG GCC CTG TTT AGG ATT AGA GAC Pro Glu Asn Ser Pro Glu Thr Glu Val Ala Leu Phe Arg Ile Arg Asp 360 365 370	1878
CGA GAC TCT GGA GAA AAT GGA AAA ATG ATT TGC TCA ATT CAG GAT GAT Arg Asp Ser Gly Glu Asn Gly Lys Met Ile Cys Ser Ile Gln Asp Asp 375 380 385	1926
GTT CCT TTT AAG CTA AAA CCT TCT GTT GAG AAT TTC TAC AGG CTG GTA Val Pro Phe Lys Leu Lys Pro Ser Val Glu Asn Phe Tyr Arg Leu Val 390 395 400	1974
ACA GAA GGG GCG CTG GAC AGA GAG ACC AGA GCC GAG TAC AAC ATC ACC Thr Glu Gly Ala Leu Asp Arg Glu Thr Arg Ala Glu Tyr Asn Ile Thr 405 410 415 420	2022
ATC ACC ATC ACA GAC TTG GGG ACT CCA AGG CTG AAA ACC GAG CAG AGC Ile Thr Ile Thr Asp Leu Gly Thr Pro Arg Leu Lys Thr Glu Gln Ser 425 430 435	2070
ATA ACC GTG CTG GTG TCG GAC GTC AAT GAC AAC GCC CCC GCC TTC ACC Ile Thr Val Leu Val Ser Asp Val Asn Asp Asn Ala Pro Ala Phe Thr 440 445 450	2118
CAA ACC TCC TAC ACC CTG TTC GTC CGC GAG AAC AAC AGC CCC GCC CTG Gln Thr Ser Tyr Thr Leu Phe Val Arg Glu Asn Asn Ser Pro Ala Leu 455 460 465	2166
CAC ATC GGC AGT GTC AGC GCC ACA GAC AGA GAC TCG GGC ACC AAC GCC His Ile Gly Ser Val Ser Ala Thr Asp Arg Asp Ser Gly Thr Asn Ala 470 475 480	2214
CAG GTC ACC TAC TCG CTG CTG CCG CCC CAG GAC CCG CAC CTG CCC CTA Gln Val Thr Tyr Ser Leu Leu Pro Pro Gln Asp Pro His Leu Pro Leu 485 490 495 500	2262
ACC TCC CTG GTC TCC ATT AAC ACG GAC AAC GGC CAC CTG TTC GCT CTC Thr Ser Leu Val Ser Ile Asn Thr Asp Asn Gly His Leu Phe Ala Leu 505 510 515	2310
CAG TCG CTG GAC TAC GAG GCC CTG CAG GCT TTC GAG TTC CGC GTG GGC Gln Ser Leu Asp Tyr Glu Ala Leu Gln Ala Phe Glu Phe Arg Val Gly 520 525 530	2358
GCC ACA GAC CGC GGC TTC CCG GCG CTG AGC AGC GAG GCG CTG GTG CGA Ala Thr Asp Arg Gly Phe Pro Ala Leu Ser Ser Glu Ala Leu Val Arg 535 540 545	2406

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Val	Leu	Val	Leu	Asp	Ala	Asn	Asp	Asn	Ser	Pro	Phe	Val	Leu	Tyr	Pro	2454
CTG	CTG	GTG	CTG	GAC	GCC	AAC	GAC	AAC	TCG	CCC	TTC	GTG	CTG	TAC	CCG	
550					555					560						
CTG	CAG	AAC	GGC	TCC	GCG	CCC	TGC	ACC	GAG	CTG	GTG	CCC	CGG	GCG	GCC	2502
Leu	Gln	Asn	Gly	Ser	Ala	Pro	Cys	Thr	Glu	Leu	Val	Pro	Arg	Ala	Ala	
565					570					575					580	
GAG	CCG	GGC	TAC	CTG	GTG	ACC	AAG	GTG	GTG	GCG	GTG	GAC	GGC	GAC	TCG	2550
Glu	Pro	Gly	Tyr	Leu	Val	Thr	Lys	Val	Val	Ala	Val	Asp	Gly	Asp	Ser	
				585					590					595		
GGC	CAG	AAC	GCC	TGG	CTG	TCG	TAC	CAG	CTG	CTC	AAG	GCC	ACG	GAG	CCC	2598
Gly	Gln	Asn	Ala	Trp	Leu	Ser	Tyr	Gln	Leu	Leu	Lys	Ala	Thr	Glu	Pro	
			600					605					610			
GGG	CTG	TTC	GGC	GTG	TGG	GCG	CAC	AAT	GGC	GAG	GTG	CGC	ACC	GCC	AGG	2646
Gly	Leu	Phe	Gly	Val	Trp	Ala	His	Asn	Gly	Glu	Val	Arg	Thr	Ala	Arg	
		615					620					625				
CTG	CTG	AGC	GAG	CGC	GAC	GTG	GCC	AAG	CAC	AGG	CTA	GTG	GTG	CTG	GTC	2694
Leu	Leu	Ser	Glu	Arg	Asp	Val	Ala	Lys	His	Arg	Leu	Val	Val	Leu	Val	
	630					635					640					
AAG	GAC	AAT	GGC	GAG	CCT	CCG	CGC	TCG	GCC	ACA	GCC	ACG	CTG	CAA	GTG	2742
Lys	Asp	Asn	Gly	Glu	Pro	Pro	Arg	Ser	Ala	Thr	Ala	Thr	Leu	Gln	Val	
645					650					655					660	
CTC	CTG	GTG	GAC	GGC	TTC	TCT	CAG	CCC	TAC	CTG	CCG	CTC	CCA	GAG	GCG	2790
Leu	Leu	Val	Asp	Gly	Phe	Ser	Gln	Pro	Tyr	Leu	Pro	Leu	Pro	Glu	Ala	
				665					670					675		
GCC	CCG	GCC	CAA	GCC	CAG	GCC	GAC	TCG	CTT	ACC	GTC	TAC	CTG	GTG	GTG	2838
Ala	Pro	Ala	Gln	Ala	Gln	Ala	Asp	Ser	Leu	Thr	Val	Tyr	Leu	Val	Val	
			680					685					690			
GCA	TTG	GCC	TCG	GTG	TCT	TCG	CTC	TTC	CTC	TTC	TCG	GTG	TTC	CTG	TTC	2886
Ala	Leu	Ala	Ser	Val	Ser	Ser	Leu	Phe	Leu	Phe	Ser	Val	Phe	Leu	Phe	
		695					700					705				
GTG	GCA	GTG	CGG	CTG	TGC	AGG	AGG	AGC	AGG	GCG	GCC	TCA	GTG	GGT	CGC	2934
Val	Ala	Val	Arg	Leu	Cys	Arg	Arg	Ser	Arg	Ala	Ala	Ser	Val	Gly	Arg	
	710					715					720					
TGC	TCG	GTG	CCC	GAG	GGC	CCC	TTT	CCA	GGG	CAT	CTG	GTG	GAC	GTG	AGC	2982
Cys	Ser	Val	Pro	Glu	Gly	Pro	Phe	Pro	Gly	His	Leu	Val	Asp	Val	Ser	
725					730					735					740	
GGC	ACC	GGG	ACC	CTT	TCC	CAG	AGC	TAC	CAG	TAC	GAG	GTG	TGT	CTG	ACG	3030
Gly	Thr	Gly	Thr	Leu	Ser	Gln	Ser	Tyr	Gln	Tyr	Glu	Val	Cys</			

TTAGTTTTTT TTAACCCTTT AGTAATCTTG AATTCTACTT TTTTTTAAAT TTCTACTGTT	3243
GTCTTTTAGTA ATGTTACTCA TTTCCTTTGT CTGATTGTTA GTTTTCAAAT TATTGTATTA	3303
TTATAAATAT TTTATATCAG GAAAGTTCAT ATTTCTGAAT AAATTAATAG	3353

(2) INFORMATION FOR SEQ ID NO:110:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 787 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:110:

Met	Glu	Pro	Ala	Gly	Glu	Arg	Phe	Pro	Glu	Gln	Arg	Gln	Val	Leu	Ile	1	5	10	15
Leu	Leu	Leu	Leu	Leu	Glu	Val	Thr	Leu	Ala	Gly	Trp	Glu	Pro	Arg	Arg	20	25	30	
Tyr	Ser	Val	Met	Glu	Glu	Thr	Glu	Arg	Gly	Ser	Phe	Val	Ala	Asn	Leu	35	40	45	
Ala	Asn	Asp	Leu	Gly	Leu	Gly	Val	Gly	Glu	Leu	Ala	Glu	Arg	Gly	Ala	50	55	60	
Arg	Val	Val	Ser	Glu	Asp	Asn	Glu	Gln	Gly	Leu	Gln	Leu	Asp	Leu	Gln	65	70	75	80
Thr	Gly	Gln	Leu	Ile	Leu	Asn	Glu	Lys	Leu	Asp	Arg	Glu	Lys	Leu	Cys	85	90	95	
Gly	Pro	Thr	Glu	Pro	Cys	Ile	Met	His	Phe	Gln	Val	Leu	Leu	Lys	Lys	100	105	110	
Pro	Leu	Glu	Val	Phe	Arg	Ala	Glu	Leu	Leu	Val	Thr	Asp	Ile	Asn	Asp	115	120	125	
His	Ser	Pro	Glu	Phe	Pro	Glu	Arg	Glu	Met	Thr	Leu	Lys	Ile	Pro	Glu	130	135	140	
Thr	Ser	Ser	Leu	Gly	Thr	Val	Phe	Pro	Leu	Lys	Lys	Ala	Arg	Asp	Leu	145	150	155	160
Asp	Val	Gly	Ser	Asn	Asn	Val	Gln	Asn	Tyr	Asn	Ile	Ser	Pro	Asn	Ser	165	170	175	
His	Phe	His	Val	Ser	Thr	Arg	Thr	Arg	Gly	Asp	Gly	Arg	Lys	Tyr	Pro	180	185	190	
Glu	Leu	Val	Leu	Asp	Thr	Glu	Leu	Asp	Arg	Glu	Glu	Gln	Ala	Glu	Leu	195	200	205	
Arg	Leu	Thr	Leu	Thr	Ala	Val	Asp	Gly	Gly	Ser	Pro	Pro	Arg	Ser	Gly	210	215	220	

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Thr Val Gln Ile Leu Ile Leu Val Leu Asp Ala Asn Asp Asn Ala Pro  
225 230 235 240

Glu Phe Val Gln Ala Leu Tyr Glu Val Gln Val Pro Glu Asn Ser Pro  
245 250 255

Val Gly Ser Leu Val Val Lys Val Ser Ala Arg Asp Leu Asp Thr Gly  
260 265 270

Thr Asn Gly Glu Ile Ser Tyr Ser Leu Tyr Tyr Ser Ser Gln Glu Ile  
275 280 285

Asp Lys Pro Phe Glu Leu Ser Ser Leu Ser Gly Glu Ile Arg Leu Ile  
290 295 300

Lys Lys Leu Asp Phe Glu Thr Met Ser Ser Tyr Asp Leu Asp Ile Glu  
305 310 315 320

Ala Ser Asp Gly Gly Gly Leu Ser Gly Lys Cys Ser Val Ser Val Lys  
325 330 335

Val Leu Asp Val Asn Asp Asn Phe Pro Glu Leu Ser Ile Ser Ser Leu  
340 345 350

Thr Ser Pro Ile Pro Glu Asn Ser Pro Glu Thr Glu Val Ala Leu Phe  
355 360 365

Arg Ile Arg Asp Arg Asp Ser Gly Glu Asn Gly Lys Met Ile Cys Ser  
370 375 380

Ile Gln Asp Asp Val Pro Phe Lys Leu Lys Pro Ser Val Glu Asn Phe  
385 390 395 400

Tyr Arg Leu Val Thr Glu Gly Ala Leu Asp Arg Glu Thr Arg Ala Glu  
405 410 415

Tyr Asn Ile Thr Ile Thr Ile Thr Asp Leu Gly Thr Pro Arg Leu Lys  
420 425 430

Thr Glu Gln Ser Ile Thr Val Leu Val Ser Asp Val Asn Asp Asn Ala  
435 440 445

Pro Ala Phe Thr Gln Thr Ser Tyr Thr Leu Phe Val Arg Glu Asn Asn  
450 455 460

Ser Pro Ala Leu His Ile Gly Ser Val Ser Ala Thr Asp Arg Asp Ser  
465 470 475 480

Gly Thr Asn Ala Gln Val Thr Tyr Ser Leu Leu Pro Pro Gln Asp Pro  
485 490 495

His Leu Pro Leu Thr Ser Leu Val Ser Ile Asn Thr Asp Asn Gly His  
500 505 510

Leu Phe Ala Leu Gln Ser Leu Asp Tyr Glu Ala Leu Gln Ala Phe Glu  
515 520 525

Phe Arg Val Gly Ala Thr Asp Arg Gly Phe Pro Ala Leu Ser Ser Glu  
530 535 540

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Ala Leu Val Arg Val Leu Val Leu Asp Ala Asn Asp Asn Ser Pro Phe  
545 550 555 560

Val Leu Tyr Pro Leu Gln Asn Gly Ser Ala Pro Cys Thr Glu Leu Val  
565 570 575

Pro Arg Ala Ala Glu Pro Gly Tyr Leu Val Thr Lys Val Val Ala Val  
580 585 590

Asp Gly Asp Ser Gly Gln Asn Ala Trp Leu Ser Tyr Gln Leu Leu Lys  
595 600 605

Ala Thr Glu Pro Gly Leu Phe Gly Val Trp Ala His Asn Gly Glu Val  
610 615 620

Arg Thr Ala Arg Leu Leu Ser Glu Arg Asp Val Ala Lys His Arg Leu  
625 630 635 640

Val Val Leu Val Lys Asp Asn Gly Glu Pro Pro Arg Ser Ala Thr Ala  
645 650 655

Thr Leu Gln Val Leu Leu Val Asp Gly Phe Ser Gln Pro Tyr Leu Pro  
660 665 670

Leu Pro Glu Ala Ala Pro Ala Gln Ala Gln Ala Asp Ser Leu Thr Val  
675 680 685

Tyr Leu Val Val Ala Leu Ala Ser Val Ser Ser Leu Phe Leu Phe Ser  
690 695 700

Val Phe Leu Phe Val Ala Val Arg Leu Cys Arg Arg Ser Arg Ala Ala  
705 710 715 720

Ser Val Gly Arg Cys Ser Val Pro Glu Gly Pro Phe Pro Gly His Leu  
725 730 735

Val Asp Val Ser Gly Thr Gly Thr Leu Ser Gln Ser Tyr Gln Tyr Glu  
740 745 750

Val Cys Leu Thr Gly Gly Ser Glu Ser Asn Asp Phe Lys Phe Leu Lys  
755 760 765

Pro Ile Phe Pro Asn Ile Val Ser Gln Asp Ser Arg Arg Lys Ser Glu  
770 775 780

Phe Leu Glu  
785

(2) INFORMATION FOR SEQ ID NO:111:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 3033 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

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(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION: 138..2528

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:111:

GTGATTGGAC GTGTTTTTGT GACTATTTGG GAAGAAGACA CCTTCCTAAT CAGATTTACT	60
CCAATATCTT CCCGGACCCT CATGAGTGGG TTGCAATTGA CTTGAAGAAG CAGCACCCTC	120
AGGACTGAAT CTGAACA ATG GAG ACA GCA CTA GCA AAA ATA CCA CAG CAA	170
Met Glu Thr Ala Leu Ala Lys Ile Pro Gln Gln	
1 5 10	
AGG CAA GTC TTT TTT CTT ACT ATA TTG TCG TTA TTG TGG AAG TCT AGC	218
Arg Gln Val Phe Phe Leu Thr Ile Leu Ser Leu Leu Trp Lys Ser Ser	
15 20 25	
TCT GAG GCC ATT AGA TAT TCC ATG CCA GAA GAA ACA GAG AGT GGC TAT	266
Ser Glu Ala Ile Arg Tyr Ser Met Pro Glu Glu Thr Glu Ser Gly Tyr	
30 35 40	
ATG GTG GCT AAC CTG GCG AAA GAT CTG GGG ATC AGG GTT GGA GAA CTG	314
Met Val Ala Asn Leu Ala Lys Asp Leu Gly Ile Arg Val Gly Glu Leu	
45 50 55	
TCC TCT AGA GGA GCT CAA ATC CAT TAC AAA GGA AAC AAA GAA CTT TTG	362
Ser Ser Arg Gly Ala Gln Ile His Tyr Lys Gly Asn Lys Glu Leu Leu	
60 65 70 75	
CAG CTG GAT GCA GAG ACT GGG AAT TTG TTC TTA AAG GAA AAA CTA GAC	410
Gln Leu Asp Ala Glu Thr Gly Asn Leu Phe Leu Lys Glu Lys Leu Asp	
80 85 90	
AGA GAA CTG CTG TGT GGA GAG ACA GAA CCC TGT GTG CTG AAC TTC CAG	458
Arg Glu Leu Leu Cys Gly Glu Thr Glu Pro Cys Val Leu Asn Phe Gln	
95 100 105	
ATC ATA CTG GAA AAC CCT ATG CAG TTC TTC CAA ACT GAA CTG CAG CTC	506
Ile Ile Leu Glu Asn Pro Met Gln Phe Phe Gln Thr Glu Leu Gln Leu	
110 115 120	
ACA GAT ATA AAC GAC CAT TCT CCA GAG TTC CCC AAC AAG AAA ATG CTT	554
Thr Asp Ile Asn Asp His Ser Pro Glu Phe Pro Asn Lys Lys Met Leu	
125 130 135	
CTA ACA ATT CCT GAG AGT GCC CAT CCA GGG ACT GTG TTT CCT CTG AAG	602
Leu Thr Ile Pro Glu Ser Ala His Pro Gly Thr Val Phe Pro Leu Lys	
140 145 150 155	
GCA GCT CGG GAC TCT GAC ATA GGG AGC AAC GCT GTT CAG AAC TAC ACA	650
Ala Ala Arg Asp Ser Asp Ile Gly Ser Asn Ala Val Gln Asn Tyr Thr	
160 165 170	
GTC AAT CCC AAC CTC CAT TTC CAC GTC GTT ACT CAC AGT CGC ACA GAT	698
Val Asn Pro Asn Leu His Phe His Val Val Thr His Ser Arg Thr Asp	
175 180 185	

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GGC AGG AAA TAC CCA GAG CTG GTG CTG GAC AGA GCC CTG GAT AGG GAG Gly Arg Lys Tyr Pro Glu Leu Val Leu Asp Arg Ala Leu Asp Arg Glu 190 195 200	746
GAG CAG CCT GAG CTC ACT TTA ATC CTC ACT GCT CTG GAT GGT GGA GCT Glu Gln Pro Glu Leu Thr Thr Ile Leu Thr Ala Leu Asp Gly Gly Ala 205 210 215	794
CCT TCC AGG TCA GGA ACC ACC ACA GTT CAC ATA GAA GTT GTG GAC ATC Pro Ser Arg Ser Gly Thr Thr Thr Val His Ile Glu Val Val Asp Ile 220 225 230 235	842
AAT GAT AAC TCC CCC CAG TTT GTA CAG TCA CTC TAT AAG GTG CAA GTT Asn Asp Asn Ser Pro Gln Phe Val Gln Ser Leu Tyr Lys Val Gln Val 240 245 250	890
CCT GAG AAT AAT CCC CTC AAT GCC TTT GTT GTC ACG GTC TCT GCC ACG Pro Glu Asn Asn Pro Leu Asn Ala Phe Val Val Thr Val Ser Ala Thr 255 260 265	938
GAT TTA GAT GCT GGG GTA TAT GGC AAT GTG ACC TAT TCT CTG TTT CAA Asp Leu Asp Ala Gly Val Tyr Gly Asn Val Thr Tyr Ser Leu Phe Gln 270 275 280	986
GGG TAT GGG GTA TTT CAA CCA TTT GTA ATA GAC GAA ATC ACT GGA GAA Gly Tyr Gly Val Phe Gln Pro Phe Val Ile Asp Glu Ile Thr Gly Glu 285 290 295	1034
ATC CAT CTG AGC AAA GAG CTG GAT TTT GAG GAA ATT AGC AAT CAT AAC Ile His Leu Ser Lys Glu Leu Asp Phe Glu Glu Ile Ser Asn His Asn 300 305 310 315	1082
ATA GAA ATC GCA GCC ACA GAT GGA GGA GGC CTT TCA GGA AAA TGC ACT Ile Glu Ile Ala Ala Thr Asp Gly Gly Gly Leu Ser Gly Lys Cys Thr 320 325 330	1130
GTG GCT GTA CAG GTG TTG GAT GTG AAT GAC AAC GCC CCA GAG TTG ACA Val Ala Val Gln Val Leu Asp Val Asn Asp Asn Ala Pro Glu Leu Thr 335 340 345	1178
ATT AGG AAG CTC ACA GTC CTG GTC CCA GAA AAT TCC CCA GAG ACT GTA Ile Arg Lys Leu Thr Val Leu Val Pro Glu Asn Ser Ala Glu Thr Val 350 355 360	1226
GTT GCT GTT TTT AGT GTT TCT GAT TCT GAT TCG GGG GAC AAT GGA AGG Val Ala Val Phe Ser Val Ser Asp Ser Asp Ser Gly Asp Asn Gly Arg 365 370 375	1274
ATG GTG TGT TCT ATT CCG AAC AAT ATC CCA TTT CTC CTG AAA CCC ACA Met Val Cys Ser Ile Pro Asn Asn Ile Pro Phe Leu Leu Lys Pro Thr 380 385 390 395	1322
TTT GAG AAT TAT TAC ACG TTA GTG ACT GAG GGG CCA CTT GAT AGA GAG Phe Glu Asn Tyr Tyr Thr Leu Val Thr Glu Gly Pro Leu Asp Arg Glu 400 405 410	1370
AAC AGA GCT GAG TAC AAC ATC ACC ATC ACG GTC TCA GAT CTG GGC ACA Asn Arg Ala Glu Tyr Asn Ile Thr Ile Thr Val Ser Asp Leu Gly Thr 415 420 425	1418

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CCC AGG CTC ACA ACC CAG CAC ACC ATA ACA GTG CAA GTG TCC GAC ATC Pro Arg Leu Thr Thr Gln His Thr Ile Thr Val Gln Val Ser Asp Ile 430 435 440	1466
AAC GAC AAC GCC CCT GCC TTC ACC CAA ACC TCC TAC ACC ATG TTT GTC Asn Asp Asn Ala Pro Ala Phe Thr Gln Thr Ser Tyr Thr Met Phe Val 445 450 455	1514
CAC GAG AAC AAC AGC CCC GCC CTG CAC ATA GGC ACC ATC AGT GCC ACA His Glu Asn Asn Ser Pro Ala Leu His Ile Gly Thr Ile Ser Ala Thr 460 465 470 475	1562
GAC TCA GAC TCA GGC TCC AAT GCC CAC ATC ACC TAC TCG CTG CTG CCG Asp Ser Asp Ser Gly Ser Asn Ala His Ile Thr Tyr Ser Leu Leu Pro 480 485 490	1610
CCT GAT GAC CCG CAG CTG GCC CTC GAC TCA CTC ATC TCC ATC AAT GTT Pro Asp Asp Pro Gln Leu Ala Leu Asp Ser Leu Ile Ser Ile Asn Val 495 500 505	1658
GAC AAT GGG CAG CTG TTC GCG CTC AGA GCT CTA GAC TAT GAG GCA CTG Asp Asn Gly Gln Leu Phe Ala Leu Arg Ala Leu Asp Tyr Glu Ala Leu 510 515 520	1706
CAG TCC TTC GAG TTC TAC GTG GGC GCT ACA GAT GGA GGC TCA CCC GCG Gln Ser Phe Glu Phe Tyr Val Gly Ala Thr Asp Gly Gly Ser Pro Ala 525 530 535	1754
CTC AGC AGC CAG ACT CTG GTG CGG ATG GTG GTG CTG GAT GAC AAT GAC Leu Ser Ser Gln Thr Leu Val Arg Met Val Val Leu Asp Asp Asn Asp 540 545 550 555	1802
AAT GCC CCC TTC GTG CTC TAC CCA CTG CAG AAT GCC TCA GCA CCC TGT Asn Ala Pro Phe Val Leu Tyr Pro Leu Gln Asn Ala Ser Ala Pro Cys 560 565 570	1850
ACT GAG CTA CTG CCT AGG GCA GCA GAG CCC GGC TAC CTG ATC ACC AAA Thr Glu Leu Leu Pro Arg Ala Ala Glu Pro Gly Tyr Leu Ile Thr Lys 575 580 585	1898
GTG GTG GCT GTG GAT CGC GAC TCT GGA CAG AAT GCT TGG CTG TCG TTC Val Val Ala Val Asp Arg Asp Ser Gly Gln Asn Ala Trp Leu Ser Phe 590 595 600	1946
CAG CTA CTT AAA GCT ACA GAG CCA GGG CTG TTC AGT GTA TGG GCA CAC Gln Leu Leu Lys Ala Thr Glu Pro Gly Leu Phe Ser Val Trp Ala His 605 610 615	1994
AAT GGT GAA GTG CGC ACC ACT AGG CTG CTG AGT GAG CGA GAT GCT CAG Asn Gly Glu Val Arg Thr Thr Arg Leu Leu Ser Glu Arg Asp Ala Gln 620 625 630 635	2042
AAG CAC AAG CTA CTG CTG GTC AAG GAC AAT GGC GAT CCT CTG CGC Lys His Lys Leu Leu Leu Val Lys Asp Asn Gly Asp Pro Leu Arg 640 645 650	2090
TCT GCC AAT GTC ACT CTT CAC GTG CTA GTG GTG GAT GGC TTC TCG CAG Ser Ala Asn Val Thr Leu His Val Leu Val Val Asp Gly Phe Ser Gln 655 660 665	2138

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CCT TAC CTA CCA TTG GCT GAG GTG GCA CAG GAT TCC ATG CAA GAT AAT Pro Tyr Leu Pro Leu Ala Glu Val Ala Gln Asp Ser Met Gln Asp Asn 670 675 680	2186
TAC GAC GTT CTC ACA CTG TAC CTA GTC ATT GCC TTG GCA TCT GTA TCT Tyr Asp Val Leu Thr Leu Tyr Leu Val Ile Ala Leu Ala Ser Val Ser 685 690 695	2234
TCT CTC TTC CTC TTG TCT GTA GTG CTG TTT GTG GGG GTG AGG CTG TGC Ser Leu Phe Leu Leu Ser Val Val Leu Phe Val Gly Val Arg Leu Cys 700 705 710 715	2282
AGG AGG GCC AGG GAG GCC TCC TTG GGT GAC TAC TCT GTG CCT GAG GGA Arg Arg Ala Arg Glu Ala Ser Leu Gly Asp Tyr Ser Val Pro Glu Gly 720 725 730	2330
CAC TTT CCT AGC CAC TTG GTG GAT GTC AGC GGT GCC GGG ACC CTG TCC His Phe Pro Ser His Leu Val Asp Val Ser Gly Ala Gly Thr Leu Ser 735 740 745	2378
CAG AGT TAT CAA TAT GAG GTG TGT CTT AAT GGA GGT ACT AGA ACA AAT Gln Ser Tyr Gln Tyr Glu Val Cys Leu Asn Gly Gly Thr Arg Thr Asn 750 755 760	2426
GAG TTT AAC TTT CTT AAA CCA TTG TTT CCT ATC CTT CCG ACC CAG GCT Glu Phe Asn Phe Leu Lys Pro Leu Phe Pro Ile Leu Pro Thr Gln Ala 765 770 775	2474
GCT GCT GCT GAA GAA AGA GAA AAC GCT GTT GTG CAC AAT AGC GTT GGA Ala Ala Ala Glu Glu Arg Glu Asn Ala Val His Asn Ser Val Gly 780 785 790 795	2522
TTC TAT TAGAGCACTG ATTTTGAAGT GGTGGTTACC TCATTTTTC TTAACATATCC Phe Tyr	2578
CTGATGTAGA ATGGTGTAGT GCCGTGAATC AACTCCTGAG ATATATGTTT ATTTTATCCT	2638
TTGTTTTGAA TCAAACATATT CAGATGTGAT CCTACTCTAG AGAATTTGGT TCTACTCCAT	2698
TGTGTTTGTGTT TAGATTTCTA CGCCATACCA GTGCATGCTG GGTGTTTTTT TTTTTTACAA	2758
TTATTATAAC TTTGCTTTGG AGGGGAACTC ATATTCGCTG TAACGAATTG GAACCACTTT	2818
CATTGTTAGA GATGCCTTGC TTTGTTGTGT TATTTTCAGAC AGGGTCTTAA ATTGTAGCCC	2878
TGGGTGACCT GAAATGACTA TGTACAGACT GACTTTGAAT TTGTGGCAGT CCATCTGCCT	2938
CTGTTGTCCT ATGTTGGGAT TGTGAGCATG CATGAGTAGG CTCAGCTGTG GTGAGCGACC	2998
TTAATAAAAA TCAAATACTA AAAAAAAAAA AAAAA	3033

(2) INFORMATION FOR SEQ ID NO:112:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 797 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:112:

Met Glu Thr Ala Leu Ala Lys Ile Pro Gln Gln Arg Gln Val Phe Phe  
1 5 10 15  
Leu Thr Ile Leu Ser Leu Leu Trp Lys Ser Ser Ser Glu Ala Ile Arg  
20 25 30  
Tyr Ser Met Pro Glu Glu Thr Glu Ser Gly Tyr Met Val Ala Asn Leu  
35 40 45  
Ala Lys Asp Leu Gly Ile Arg Val Gly Glu Leu Ser Ser Arg Gly Ala  
50 55 60  
Gln Ile His Tyr Lys Gly Asn Lys Glu Leu Leu Gln Leu Asp Ala Glu  
65 70 75 80  
Thr Gly Asn Leu Phe Leu Lys Glu Lys Leu Asp Arg Glu Leu Leu Cys  
85 90 95  
Gly Glu Thr Glu Pro Cys Val Leu Asn Phe Gln Ile Ile Leu Glu Asn  
100 105 110  
Pro Met Gln Phe Phe Gln Thr Glu Leu Gln Leu Thr Asp Ile Asn Asp  
115 120 125  
His Ser Pro Glu Phe Pro Asn Lys Lys Met Leu Leu Thr Ile Pro Glu  
130 135 140  
Ser Ala His Pro Gly Thr Val Phe Pro Leu Lys Ala Ala Arg Asp Ser  
145 150 155 160  
Asp Ile Gly Ser Asn Ala Val Gln Asn Tyr Thr Val Asn Pro Asn Leu  
165 170 175  
His Phe His Val Val Thr His Ser Arg Thr Asp Gly Arg Lys Tyr Pro  
180 185 190  
Glu Leu Val Leu Asp Arg Ala Leu Asp Arg Glu Glu Gln Pro Glu Leu  
195 200 205  
Thr Leu Ile Leu Thr Ala Leu Asp Gly Gly Ala Pro Ser Arg Ser Gly  
210 215 220  
Thr Thr Thr Val His Ile Glu Val Val Asp Ile Asn Asp Asn Ser Pro  
225 230 235 240  
Gln Phe Val Gln Ser Leu Tyr Lys Val Gln Val Pro Glu Asn Asn Pro  
245 250 255  
Leu Asn Ala Phe Val Val Thr Val Ser Ala Thr Asp Leu Asp Ala Gly  
260 265 270  
Val Tyr Gly Asn Val Thr Tyr Ser Leu Phe Gln Gly Tyr Gly Val Phe  
275 280 285  
Gln Pro Phe Val Ile Asp Glu Ile Thr Gly Glu Ile His Leu Ser Lys  
290 295 300

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Glu Leu Asp Phe Glu Glu Ile Ser Asn His Asn Ile Glu Ile Ala Ala  
 305 310 315 320  
 Thr Asp Gly Gly Gly Leu Ser Gly Lys Cys Thr Val Ala Val Gln Val  
 325 330 335  
 Leu Asp Val Asn Asp Asn Ala Pro Glu Leu Thr Ile Arg Lys Leu Thr  
 340 345 350  
 Val Leu Val Pro Glu Asn Ser Ala Glu Thr Val Val Ala Val Phe Ser  
 355 360 365  
 Val Ser Asp Ser Asp Ser Gly Asp Asn Gly Arg Met Val Cys Ser Ile  
 370 375 380  
 Pro Asn Asn Ile Pro Phe Leu Leu Lys Pro Thr Phe Glu Asn Tyr Tyr  
 385 390 395 400  
 Thr Leu Val Thr Glu Gly Pro Leu Asp Arg Glu Asn Arg Ala Glu Tyr  
 405 410 415  
 Asn Ile Thr Ile Thr Val Ser Asp Leu Gly Thr Pro Arg Leu Thr Thr  
 420 425 430  
 Gln His Thr Ile Thr Val Gln Val Ser Asp Ile Asn Asp Asn Ala Pro  
 435 440 445  
 Ala Phe Thr Gln Thr Ser Tyr Thr Met Phe Val His Glu Asn Asn Ser  
 450 455 460  
 Pro Ala Leu His Ile Gly Thr Ile Ser Ala Thr Asp Ser Asp Ser Gly  
 465 470 475 480  
 Ser Asn Ala His Ile Thr Tyr Ser Leu Leu Pro Pro Asp Asp Pro Gln  
 485 490 495  
 Leu Ala Leu Asp Ser Leu Ile Ser Ile Asn Val Asp Asn Gly Gln Leu  
 500 505 510  
 Phe Ala Leu Arg Ala Leu Asp Tyr Glu Ala Leu Gln Ser Phe Glu Phe  
 515 520 525  
 Tyr Val Gly Ala Thr Asp Gly Gly Ser Pro Ala Leu Ser Ser Gln Thr  
 530 535 540  
 Leu Val Arg Met Val Val Leu Asp Asp Asn Asp Asn Ala Pro Phe Val  
 545 550 555 560  
 Leu Tyr Pro Leu Gln Asn Ala Ser Ala Pro Cys Thr Glu Leu Leu Pro  
 565 570 575  
 Arg Ala Ala Glu Pro Gly Tyr Leu Ile Thr Lys Val Val Ala Val Asp  
 580 585 590  
 Arg Asp Ser Gly Gln Asn Ala Trp Leu Ser Phe Gln Leu Leu Lys Ala  
 595 600 605  
 Thr Glu Pro Gly Leu Phe Ser Val Trp Ala His Asn Gly Glu Val Arg  
 610 615 620

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Thr Thr Arg Leu Leu Ser Glu Arg Asp Ala Gln Lys His Lys Leu Leu  
625 630 635 640

Leu Leu Val Lys Asp Asn Gly Asp Pro Leu Arg Ser Ala Asn Val Thr  
645 650 655

Leu His Val Leu Val Val Asp Gly Phe Ser Gln Pro Tyr Leu Pro Leu  
660 665 670

Ala Glu Val Ala Gln Asp Ser Met Gln Asp Asn Tyr Asp Val Leu Thr  
675 680 685

Leu Tyr Leu Val Ile Ala Leu Ala Ser Val Ser Ser Leu Phe Leu Leu  
690 695 700

Ser Val Val Leu Phe Val Gly Val Arg Leu Cys Arg Arg Ala Arg Glu  
705 710 715 720

Ala Ser Leu Gly Asp Tyr Ser Val Pro Glu Gly His Phe Pro Ser His  
725 730 735

Leu Val Asp Val Ser Gly Ala Gly Thr Leu Ser Gln Ser Tyr Gln Tyr  
740 745 750

Glu Val Cys Leu Asn Gly Gly Thr Arg Thr Asn Glu Phe Asn Phe Leu  
755 760 765

Lys Pro Leu Phe Pro Ile Leu Pro Thr Gln Ala Ala Ala Ala Glu Glu  
770 775 780

Arg Glu Asn Ala Val Val His Asn Ser Val Gly Phe Tyr  
785 790 795

(2) INFORMATION FOR SEQ ID NO:113:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 2347 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:113:

AAAACACGGG GGAAATGACA GTAGCAAAGA ATCTGGACTA TGAAGAATGC TCATTGTATG	60
AAATGGAAAT ACAGGCTGAA GATGTGGGGG CGCTTCTGGG GAGGAGCAAA GTGGTAATTA	120
TGGTAGAAGA TGTAATGAC AATCGGCCAG AAGTGACCAT TACATCCTTG TTTAACCCGG	180
TATTGGAAAA TTCTCTTCCC GGGACAGTAA TTGCCTTCTT GAATGTGCAT GACCGAGACT	240
CTGGAAAGAA CGGCCAAGTT GTCTGTTACA CGCATGATAA CTTACCTTTT AAATTAGAAA	300
AGTCAATAGA TAATTATTAT AGATTGGTGA CATGGAAATA TTTGGACCGA GAAAAAGTCT	360
CCATCTACAA TATCACAGTG ATAGCCTCAG ATCTAGGAGC CCACTCTGTC ACTGAAACTT	420



ACATTGCCCT GATTGTGGCA GACACTAATG ACAACCCTCC TCGTTTTCCT CACACCTCCT	480
ACACAGCCTA TATTCCAGAG AACAACTGA GGGGCGCCTC CATCTTCTCA CTGACTGCAC	540
ATGATCCTGA CAGTCAGGAA AATGCACAGG TCACTTACTC TGTGTCTGAG GACACCATAC	600
AGGGAGTGCC TTTGTCTCT TATATCTCCA TCAACTCAGA TACTGGTGTC CTGTATGCAC	660
TGCACTCTTT TGACTTCGAG AAGATACAAG ACTTGCAGCT ACTGGTTGTT GCCACTGACA	720
GTGGAAGCCC ACCTCTCAGC AGCAATGTGT CATTGAGCTT GTTTGTGTTG GACCAGAACG	780
ACAACGCACC TGAGATTCTA TATCCTAGCT TCCCCACAGA TGGCTCCACT GGTGTGGAAC	840
TAGCACCCCG CTCTGCAGAG CCTGGATACC TAGTGACCAA AGTGGTGGCA GTGGACAAAG	900
ACTCAGGACA GAATGCTTGG CTGTCCTACC GTCTGCTGAA GGCCAGCGAA CCTGGGCTCT	960
TCTCTGTAGG ACTTCACACG GGTGAGGTGC GTACAGCGAG GGCCCTGCTG GACAGAGATG	1020
CTCTCAAACA GAATCTGGTG ATGGCCGTGC AGGACCATGG CCAACCCCT CTCTCGGCCA	1080
CTGTAACCTCT CACTGTGGCA GTGGCTAACA GCATCCCTGA GGTGTTGGCT GACTTGAGCA	1140
GCATTAGGAC CCCTGGGGTA CCAGAGGATT CTGATATCAC GCTCCACCTG GTGGTGGCAG	1200
TGGCTGTGGT CTCCTGTGTC TTCCTTGTCT TTGTCATTGT CCTCCTAGCT CTCAGGCTTC	1260
AGCGCTGGCA GAAGTCTCGC CAGCTCCAGG GCTCCAAAGG TGGATTGGCT CCTGCACCTC	1320
CATCACATTT TGTGGGCATC GACGGGGTAC AGGCTTTTCT ACAAACCTAT TCTCATGAAG	1380
TCTCGCTCAC TTCAGGCTCC CAGACAAGCC ACATTATCTT TCCTCAGCCC AACTATGCAG	1440
ACATGCTCAT TAACCAAGAA GGCTGTGAGA AAAATGATTC CTTATTAACA TCCATAGATT	1500
TTCATGAGAG TAACCGTGAA GATGCTTGCG CCCCAGCAAGC CCCGCCAAC ACTGACTGGC	1560
GTTTCTCTCA AGCCCAGAGA CCCGGCACGA GCGGATCCCA AAATGGGGAT GAAACCGGCA	1620
CCTGGCCCAA CAACCAGTTC GATACAGAGA TGCTGCAAGC CATGATCTTG GCCTCTGCCA	1680
GTGAAGCCGC TGATGGGAGC TCCACTCTGG GAGGGGGCAC TGGCACTATG GGTTTGAGCG	1740
CTCGATATGG ACCCCAGTTT ACCCTGCAGC ACGTGCCTGA CTACCGCCAG AACGTGTACA	1800
TCCCTGGCAG CAATGCCACA CTGACCAACG CAGCTGGCAA ACGAGATGGC AAGGCTCCGG	1860
CAGGCGGCAA TGGCAACAAC AACAACTCGG GCAAGAAAGA GAAGAAGTAA TATGGAGGCC	1920
AGGCCTTGAG CCACAGGGCA GCCTCCCTCC CCAGCCAGTC CAGCTGTCC TTAATTGTAC	1980
CCAGGCCTCA GAATTTTCAAG GCTCACCCCA GGATTCTGGT AGGAGCCACA GCCAGGCCAT	2040
GCTCCCGTT GGGAAACAGA AACAAAGTGC CAAGCCAACA CCCCCTCTTT GTACCCTAGG	2100
GGGTTGAAT ATGCAAAGAG AGTTCTGCTG GGACCCCTA TCCAATCAGT GATTGTACCC	2160
ACATAGGTAG CAGGGTTAGT GTGGATACAC ACACACACAC ACACACACAC ACACACACAA	2220
CCCTTGTCTT CCGCAGTGCC TGCCACTTTC TGGGACTTTC TCATCCCCCT ACGCCCTTCC	2280

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2340

2347

(A) LENGTH: 2972 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: single  
(D) TOPOLOGY: linear

(ix) **FEATURE:**

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:114:

46

94

142

190

238

286

334

382

430

478

ACA GTA ATT GCC TTC TTG AGT GTG CAT GAC CAA GAC TCT GGA AAG AAT Thr Val Ile Ala Phe Leu Ser Val His Asp Gln Asp Ser Gly Lys Asn 160 165 170 175	526
GGT CAA GTT GTC TGT TAC ACA CGT GAT AAT TTA CCT TTT AAA TTA GAA Gly Gln Val Val Cys Tyr Thr Arg Asp Asn Leu Pro Phe Lys Leu Glu 180 185 190	574
AAG TCA ATA GGT AAT TAT TAT AGA TTA GTG ACA AGG AAA TAT TTG GAC Lys Ser Ile Gly Asn Tyr Tyr Arg Leu Val Thr Arg Lys Tyr Leu Asp 195 200 205	622
CGA GAA AAT GTC TCT ATC TAC AAT ATC ACA GTG ATG GCC TCA GAT CTA Arg Glu Asn Val Ser Ile Tyr Asn Ile Thr Val Met Ala Ser Asp Leu 210 215 220	670
GGA ACA CCA CCT CTG TCC ACT GAA ACT CAA ATC GCT CTG CAC GTG GCA Gly Thr Pro Pro Leu Ser Thr Glu Thr Gln Ile Ala Leu His Val Ala 225 230 235	718
GAC ATT AAC GAC AAC CCT CCT ACT TTC CCT CAT GCC TCC TAC TCA GCG Asp Ile Asn Asp Asn Pro Pro Thr Phe Pro His Ala Ser Tyr Ser Ala 240 245 250 255	766
TAT ATC CTA GAG AAC AAC CTG AGA GGA GCC TCC ATC TTT TCC TTG ACT Tyr Ile Leu Glu Asn Asn Leu Arg Gly Ala Ser Ile Phe Ser Leu Thr 260 265 270	814
GCA CAC GAC CCC GAC AGC CAG GAG AAT GCC CAG GTC ACT TAC TCT GTG Ala His Asp Pro Asp Ser Gln Glu Asn Ala Gln Val Thr Tyr Ser Val 275 280 285	862
ACC GAG GAC ACG CTG CAG GGG GCG CCC CTG TCC TCG TAT ATC TCC ATC Thr Glu Asp Thr Leu Gln Gly Ala Pro Leu Ser Ser Tyr Ile Ser Ile 290 295 300	910
AAC TCT GAC ACC GGT GTC CTG TAT GCG CTG CAA TCT TTC GAC TAT GAG Asn Ser Asp Thr Gly Val Leu Tyr Ala Leu Gln Ser Phe Asp Tyr Glu 305 310 315	958
CAG ATC CGA GAC CTG CAG CTA CTG GTA ACA GCC AGC GAC AGC GGG GAC Gln Ile Arg Asp Leu Gln Leu Leu Val Thr Ala Ser Asp Ser Gly Asp 320 325 330 335	1006
CCG CCC CTC AGC AGC AAC ATG TCA CTG AGC CTG TTC GTG CTG GAC CAG Pro Pro Leu Ser Ser Asn Met Ser Leu Ser Leu Phe Val Leu Asp Gln 340 345 350	1054
AAT GAC AAC GCG CCC GAG ATC CTG TAC CCC GCC CTC CCC ACA GAC GGT Asn Asp Asn Ala Pro Glu Ile Leu Tyr Pro Ala Leu Pro Thr Asp Gly 355 360 365	1102
TCC ACT GGC GTG GAG CTG GCG CCC CGC TCC GCA GAG CGT GGC TAC CTG Ser Thr Gly Val Glu Leu Ala Pro Arg Ser Ala Glu Arg Gly Tyr Leu 370 375 380	1150
GTG ACC AAG GTG GTG GCG GTG GAC AGA GAC TCG GGC CAG AAC GCC TGG Val Thr Lys Val Val Ala Val Asp Arg Asp Ser Gly Gln Asn Ala Trp 385 390 395	1198

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CTG TCC TAC CGC CTG CTC AAG GCC AGC GAG CCG GGA CTC TTC TCG GTG Leu Ser Tyr Arg Leu Leu Lys Ala Ser Glu Pro Gly Leu Phe Ser Val 400 405 410 415	1246
GGT CTG CAC ACG GGC GAG GTG CGC ACG GCG CGA GCC CTG CTG GAC AGA Gly Leu His Thr Gly Glu Val Arg Thr Ala Arg Ala Leu Leu Asp Arg 420 425 430	1294
GAC GCG CTC AAG CAG AGC CTC GTG GTG GCC GTC CAG GAC CAT GGC CAG Asp Ala Leu Lys Gln Ser Leu Val Val Ala Val Gln Asp His Gly Gln 435 440 445	1342
CCC CCT CTC TCC GCC ACT GTC ACG CTC ACC GTA GCC GTG GCT GAC AGC Pro Pro Leu Ser Ala Thr Val Thr Leu Thr Val Ala Val Ala Asp Ser 450 455 460	1390
ATC CCC GAA GTC CTG ACC GAG TTG GGC AGT CTG AAG CCT TCG GTC GAC Ile Pro Glu Val Leu Thr Glu Leu Gly Ser Leu Lys Pro Ser Val Asp 465 470 475	1438
CCG AAC GAT TCG AGC CTT ACA CTC TAT CTC GTG GTG GCA GTG GCT GCC Pro Asn Asp Ser Ser Leu Thr Leu Tyr Leu Val Val Ala Val Ala Ala 480 485 490 495	1486
ATC TCC TGT GTC TTC CTC GCC TTT GTC GCT GTG CTT CTG GGG CTC AGG Ile Ser Cys Val Phe Leu Ala Phe Val Ala Val Leu Leu Gly Leu Arg 500 505 510	1534
CTG AGG CGC TGG CAC AAG TCA CGC CTG CTC CAG GAT TCC GGT GGC AGA Leu Arg Arg Trp His Lys Ser Arg Leu Leu Gln Asp Ser Gly Gly Arg 515 520 525	1582
TTG GTA GGC GTG CCT GCC TCA CAT TTT GTG GGT GTT GAG GAG GTA CAG Leu Val Gly Val Pro Ala Ser His Phe Val Gly Val Glu Glu Val Gln 530 535 540	1630
GCT TTC CTG CAG ACC TAT TCC CAG GAA GTC TCC CTC ACC GCC GAC TCG Ala Phe Leu Gln Thr Tyr Ser Gln Glu Val Ser Leu Thr Ala Asp Ser 545 550 555	1678
CGG AAG AGT CAC CTG ATC TTT CCC CAG CCC AAC TAC GCA GAC ATG CTC Arg Lys Ser His Leu Ile Phe Pro Gln Pro Asn Tyr Ala Asp Met Leu 560 565 570 575	1726
ATC AGT CAG GAG GGC TGT GAG AAA AAT GAT TCT TTG TTA ACA TCC GTA Ile Ser Gln Glu Gly Cys Glu Lys Asn Asp Ser Leu Leu Thr Ser Val 580 585 590	1774
GAT TTT CAT GAA TAT AAG AAT GAA GCT GAT CAT GGT CAG GTG AGT TTA Asp Phe His Glu Tyr Lys Asn Glu Ala Asp His Gly Gln Val Ser Leu 595 600 605	1822
GTT CTT TGC TTG CTT TTA ATT TCC AGA TGAATTTTAT TTGGCATAAA Val Leu Cys Leu Leu Leu Ile Ser Arg 610 615	1869
TTATGTTTTG AAAACATTG TGAAGATAGT TGAAATAAT TTTTAAGGTG TATCACAGAG	1929
TTTTGGGTTT ATTTTGGTGG TGTTACCAA AAATTGAACT CTAATAGTCA TAGGTTATTG	1989
TTTCATTTGC TTTTAAACGA CTTGAAAAAG ATTGTTCCAC CATTTTAAAC CTTCCAGTAT	2049

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TTTATTCTTA TTATCACTCA TTCACTTAAG AAGTAGCTAC CCGTCCATAC TGGTAATTTT 2109  
GCTATTGTTT GTTTGTGTGT GTGTGTGTGT GTGTGTGTGT GTGTGTGTAT CCCAACTAG 2169  
AACTTCAGAA AATTATCAAG AAGTCTAAAG CCTTGTTATT AGCTTAGCAA AAGTAAATA 2229  
TATCTCAGAA TTTTAGGGT TATGTTTAGC ATTTGAACCT GTAAGTAGGC TCTTGATAT 2289  
TTCTTCACTT TAAACCTCTT TTCTGAGCCC TGTTCCTGTA CCAGTGCCCT TCAAACTTT 2349  
AATACTTCTT ACCATCCTTC AAAACATGAA CAACTTTAA AGATGGATCT TGGTGGGAGA 2409  
TGAGACTGGT TACTAAATAT TAAGTATGTG AGTCAGTGGT CACCTGGGCT CCATCCCCAT 2469  
GGAGACATGA AATCTAAAGC CTAGAATGTC CATTGCTCCC CCAAACAAAA AACAAAAGCA 2529  
AAAACATTAG ATCTGAATTA AAATGTAATT TTAACTGTT GAAAGTGAAT TTTGTAAAT 2589  
ATGTAAGAAC ATATTTCAAT ACAATTCCAA TTAGCTGTTT CGGTTGTGCA TTGATGTGAA 2649  
GTGGTGAGAA TGTGATATT AAGAACCAAT GTTTCAGGTA CACAAGTTCT AAATAAGCTG 2709  
ATCAATTCAA TTAAAGTTAT TCAGTCTTGG CTGGACACAG TGCCTCATGT CTGAAATCCC 2769  
AGCACTTTGG GAGGCTGGGG CAGGAGGACC GCTTGAGCCC CGGGGGTTTG AACTGCACT 2829  
GAGCTATGAT CATGCCACTG CACTCCAGCC TAGGTGGCAG AACTAGACCC TGTCTCTAAA 2889  
AAAACATTA TTAGGCCGCG TCGGTGGCT CACGCCTGTA ATCCCAGCAC TTTGGGAGAC 2949  
TGAGGTGGGT GGATCACCTG AGC 2972

(2) INFORMATION FOR SEQ ID NO:115:

- (i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 616 amino acids  
(B) TYPE: amino acid  
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:115:

Glu Ala Ala His His Leu Val Leu Thr Ala Ser Asp Gly Gly Lys Pro  
1 5 10 15  
Pro Arg Ser Ser Thr Val Arg Ile His Val Thr Val Leu Asp Thr Asn  
20 25 30  
Asp Asn Ala Pro Val Phe Pro His Pro Ile Tyr Arg Val Lys Val Leu  
35 40 45  
Glu Asn Met Pro Pro Gly Thr Arg Leu Leu Thr Val Thr Ala Ser Asp  
50 55 60  
Pro Asp Glu Gly Ile Asn Gly Lys Val Ala Tyr Lys Phe Arg Lys Ile  
65 70 75 80

Asn Glu Lys Gln Thr Pro Leu Phe Gln Leu Asn Glu Asn Thr Gly Glu  
85 90 95

Ile Ser Ile Ala Lys Ser Leu Asp Tyr Glu Glu Cys Ser Phe Tyr Glu  
100 105 110

Met Glu Ile Gln Ala Glu Asp Val Gly Ala Leu Leu Gly Arg Thr Lys  
115 120 125

Leu Leu Ile Ser Val Glu Asp Val Asn Asp Asn Arg Pro Glu Val Ile  
130 135 140

Ile Thr Ser Leu Phe Ser Pro Val Leu Glu Asn Ser Leu Pro Gly Thr  
145 150 155 160

Val Ile Ala Phe Leu Ser Val His Asp Gln Asp Ser Gly Lys Asn Gly  
165 170 175

Gln Val Val Cys Tyr Thr Arg Asp Asn Leu Pro Phe Lys Leu Glu Lys  
180 185 190

Ser Ile Gly Asn Tyr Tyr Arg Leu Val Thr Arg Lys Tyr Leu Asp Arg  
195 200 205

Glu Asn Val Ser Ile Tyr Asn Ile Thr Val Met Ala Ser Asp Leu Gly  
210 215 220

Thr Pro Pro Leu Ser Thr Glu Thr Gln Ile Ala Leu His Val Ala Asp  
225 230 235 240

Ile Asn Asp Asn Pro Pro Thr Phe Pro His Ala Ser Tyr Ser Ala Tyr  
245 250 255

Ile Leu Glu Asn Asn Leu Arg Gly Ala Ser Ile Phe Ser Leu Thr Ala  
260 265 270

His Asp Pro Asp Ser Gln Glu Asn Ala Gln Val Thr Tyr Ser Val Thr  
275 280 285

Glu Asp Thr Leu Gln Gly Ala Pro Leu Ser Ser Tyr Ile Ser Ile Asn  
290 295 300

Ser Asp Thr Gly Val Leu Tyr Ala Leu Gln Ser Phe Asp Tyr Glu Gln  
305 310 315 320

Ile Arg Asp Leu Gln Leu Leu Val Thr Ala Ser Asp Ser Gly Asp Pro  
325 330 335

Pro Leu Ser Ser Asn Met Ser Leu Ser Leu Phe Val Leu Asp Gln Asn  
340 345 350

Asp Asn Ala Pro Glu Ile Leu Tyr Pro Ala Leu Pro Thr Asp Gly Ser  
355 360 365

Thr Gly Val Glu Leu Ala Pro Arg Ser Ala Glu Arg Gly Tyr Leu Val  
370 375 380

Thr Lys Val Val Ala Val Asp Arg Asp Ser Gly Gln Asn Ala Trp Leu  
385 390 395 400

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Ser Tyr Arg Leu Leu Lys Ala Ser Glu Pro Gly Leu Phe Ser Val Gly  
405 410 415

Leu His Thr Gly Glu Val Arg Thr Ala Arg Ala Leu Leu Asp Arg Asp  
420 425 430

Ala Leu Lys Gln Ser Leu Val Val Ala Val Gln Asp His Gly Gln Pro  
435 440 445

Pro Leu Ser Ala Thr Val Thr Leu Thr Val Ala Val Ala Asp Ser Ile  
450 455 460

Pro Glu Val Leu Thr Glu Leu Gly Ser Leu Lys Pro Ser Val Asp Pro  
465 470 475 480

Asn Asp Ser Ser Leu Thr Leu Tyr Leu Val Val Ala Val Ala Ala Ile  
485 490 495

Ser Cys Val Phe Leu Ala Phe Val Ala Val Leu Leu Gly Leu Arg Leu  
500 505 510

Arg Arg Trp His Lys Ser Arg Leu Leu Gln Asp Ser Gly Gly Arg Leu  
515 520 525

Val Gly Val Pro Ala Ser His Phe Val Gly Val Glu Glu Val Gln Ala  
530 535 540

Phe Leu Gln Thr Tyr Ser Gln Glu Val Ser Leu Thr Ala Asp Ser Arg  
545 550 555 560

Lys Ser His Leu Ile Phe Pro Gln Pro Asn Tyr Ala Asp Met Leu Ile  
565 570 575

Ser Gln Glu Gly Cys Glu Lys Asn Asp Ser Leu Leu Thr Ser Val Asp  
580 585 590

Phe His Glu Tyr Lys Asn Glu Ala Asp His Gly Gln Val Ser Leu Val  
595 600 605

Leu Cys Leu Leu Leu Ile Ser Arg  
610 615

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